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(71)(72) Applicant and Inventor: REED, John, C. [US/US]; 4375 Toulomne Place, Carlsbad, CA 92008 (US).			
(74) Agent: BROTMAN, Harris, F.; Gray Cary Ware & Freidenrich, Suite 1700, 401 "B" Street, San Diego, CA 92101 (US).			
(54) Title: REGULATION OF bcl-2 GENE EXPRESSION			
(57) Abstract The present invention provides novel anticode oligomers and methods of using them for controlling the growth of cancer cells expressing the bcl-2 gene.			

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REGULATION OF bcl-2 GENE EXPRESSION

Reference to Government Grants

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10 Field of the Invention
The present invention relates to the field of treatments for cancer and more particularly to the field 10 of anticode oligomer treatments for cancer.

15 Related Application Data
This application is a continuation-in-part of Serial No. 07/840,716 filed February 21, 1992, which was a continuation in part of Serial No. 07,288,692 filed December 22, 1988, which has been abandoned.

20 Background of the Invention
Current approaches to cancer treatment suffer from a lack of specificity. The majority of drugs that have been developed are natural products or derivatives that either block enzyme pathways or randomly interact with DNA. Due to low therapeutic indices, most cancer treatment drugs are accompanied by serious dose-limiting toxicities. The administration of drugs to treat cancer kills not only cancer cells but also normal non-cancerous 25 cells. Because of these deleterious effects, treatments that are more specific for cancerous cells are needed.

30 It has been found that a class of genes, the oncogenes, plays a large role in the transformation and maintenance of the cancerous state and that turning off these genes, or otherwise inhibiting their effects, can return a cell to a normal phenotype. The role of oncogenes in the etiology of many human cancers has been

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reviewed in Bishop, "Cellular Oncogenes and Retroviruses," *Science*, 235:305-311 (1987). In many types of human tumors, including lymphomas and leukemias, the human bcl-2 gene is overexpressed, and may be associated with tumorigenicity (Tsujimoto et al. Involvement of the bcl-2 gene in human follicular lymphoma, *Science* 228:1440-1443 (1985)).

Antisense oligodeoxynucleotides are one example of a specific therapeutic tool with the potential for ablating oncogene function. These short (usually about 30 bases) single-stranded synthetic DNAs have a complementary base sequence to the target mRNA and form a hybrid duplex by hydrogen bonded base pairing. This hybridization can be expected to prevent expression of the target mRNA code into its protein product and thus preclude subsequent effects of the protein product. Because the mRNA sequence expressed by the gene is termed the sense sequence, the complementary sequence is termed the antisense sequence. Under some circumstances, inhibition of mRNA would be more efficient than inhibition of an enzyme's active site, since one mRNA molecule gives rise to multiple protein copies.

Synthetic oligodeoxynucleotides complementary to (antisense) mRNA of the c-myc oncogene have been used to specifically inhibit production of c-myc protein, thus arresting the growth of human leukemic cells in vitro, Holt et al., *Mol. Cell Biol.* 8:963-973 (1988), and Wickstrom et al., *Proc. Natl. Acad. Sci. USA*, 85:1028-1-32 (1988). Oligodeoxynucleotides have also been employed as specific inhibitors of retroviruses, including the human immunodeficiency virus (HIV-I), Zamecnik and Stephenson, *Proc. Natl. Acad. Sci. USA*, 75:280-284 (1978) and Zamecnik et al., *Proc. Natl. Acad. Sci. USA*, 83:4143-4146 (1986).

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Summary of the Invention

The invention provides anticodc oligomers and methods for inhibiting growth of cancer cells. The growth of lymphoma or leukemia cells, which are types of lymphocytes, are inhibited by the anticodc oligomers and methods of the invention. An anticodc oligomer complementary to at least an effective portion of the mRNA sense strand to the human *bcl-2* gene is provided and cells are then contacted with the anticodc oligomer in a concentration sufficient to inhibit growth of the cells. The methods of the invention are suitable for inhibiting growth of lymphoma/leukemia cells that express the human *bcl-2* gene and have a t (14; 18) chromosomal translocation as well as those that express the *bcl-2* gene but do not have a t (14; 18) chromosomal translocation.

In accordance with preferred embodiments, the anticodc oligomer is substantially complementary to a strategic site in the pre-mRNA sense strand or substantially complementary to the mRNA. A preferred strategic site is the translation-initiation site of the pre-mRNA coding strand. Alternative strategic sites include coding sites for splicing, transport or degradation. The subject anticodc oligomer either in its "native," unmodified form -- oligonucleotide -- or as a derivative, is brought into contact with the target lymphoma or leukemia cells. For in vivo therapeutic use, a derivative of the "native" oligonucleotide, such as the phosphorothioate form is preferable since it is believed that these forms are more resistant to degradation, notwithstanding the fact that response times to some analogues, such as the phosphorothioate analogs, has been found to be somewhat slower than to the "native" form of the oligonucleotide.

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A preferred anticod oligomer, denominated herein the TI-AS (translation initiation anticod oligomer) is an oligodeoxynucleotide which straddles the translation-initiation site of the mRNA coding strand of the human *bcl-2* gene and is complementary to this region. More preferably, this nucleotide comprises a TAC portion which is complementary to the ATG initiation sequence of the coding strand for the *bcl-2* gene, and preferably further comprises flanking portions of two to about one hundred bases, more preferably from about five to about twenty bases, which are complementary to portions of the *bcl-2* gene coding strand flanking said initiation sequence. The TI-AS nucleotide has been found effective at inhibiting the growth of the target cells both in the presence and absence of serum.

Alternatively, the anticod oligomer comprises an antisense nucleotide complementary to at least an effective portion of the splice donor site of the pre-mRNA coding strand for the human *bcl-2* gene. More particularly, this nucleotide comprises a CA portion which is complementary to the GT splice donor of the *bcl-2*, and again comprises flanking portions of two to about one hundred bases, preferably from about five to about twenty bases, which are complementary to portions of the *bcl-2* gene coding strand flanking said splice donor.

In yet another embodiment, the anticod oligomer is complementary to at least an effective portion of the splice acceptor region of the pre-mRNA coding strand for the human *bcl-2* gene. This oligomer comprises at least a TC portion which is complementary to the AG splice acceptor of the *bcl-2* gene, and again comprises flanking portions of two to about one hundred, preferably from about five to about twenty bases which are complementary to portions of the *bcl-2* gene coding

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strand flanking said acceptor. The subject oligomer may also be selected to overlap the coding site for the 26 kDa protein, *bcl-2-alpha* or for the 22 kDa protein, *bcl-2-beta*, protein products of the *bcl-2* gene.

5 Preferably the oligomer is selected to minimize homology with anticodc oligomers for pre-mRNA or mRNA coding strands for other gene sequences.

10 Accordingly, a primary object of the present invention is the provision of novel anticodc oligomers, which are useful in inhibiting the growth of cancer cells. The present invention also includes compositions for inhibiting the growth of tumor cells, which compositions comprise the anticodc oligomer of the present invention together with a pharmaceutically acceptable carrier.

15 A further object of the present invention is the provision of methods for inhibiting the growth of cancer cells using said anticodc oligomers. As a feature of the present invention, it was discovered that average reductions of 30-40% in the relative levels of *bcl-2* protein markedly enhanced the sensitivity of lymphoma cells, in particular, t(14;18)-containing lymphoma cell lines to cancer chemotherapeutic agents, including conventional anticancer drugs. Such reductions were 20 achieved by introducing into tumor cells an anticodc oligomer which binds to either pre-mRNA or mRNA expressed from the *bcl-2* gene. Two methods were used in the present invention to introduce said anticodc oligomers to tumor cells. One method involved contacting the tumor 25 cells with a composition comprising the anticodc oligomers. Another method involved transfecting the tumor cells with a vector encoding an antisense oligonucleotide. Introducing an anticodc oligomer to tumor cells achieved a reduction of *bcl-2* expression and 30

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increases the chemosensitivity of neoplastic cells to cancer chemotherapeutic agents or anticancer drugs.

Accordingly, the present invention achieved a method of killing tumor cells by introducing to tumor cells anticode oligomers which reduce *bcl-2* gene expression or impair *Bcl-2* protein function before contacting the cells with cancer chemotherapeutic agents. The cancer chemotherapeutic agents reduced the numbers of viable malignant cells, and the portion of tumor cells killed was greater than the portion which would have been killed by the same amount of drug in the absence of introducing the anticode oligomer oligodeoxynucleotide to the cells.

These and other objects of the present invention will become apparent from the following detailed description.

Brief Description of the Drawings

Figures 1 (a - d) show graphs of the effects of varying concentrations of antisense oligodeoxynucleotides on inhibition of cell proliferation.

Figures 2(a) and (b) show graphs of the concentration dependence of inhibition of cell proliferation by antisense normal and phosphorothioate oligodeoxynucleotides. Oligodeoxynucleotide additions to cultures included TI-AS phosphorothioate (○ and △; two separate experiments), TI-S phosphorothioate (Δ), TI-AS normal (□), and TI-S normal (△).

Figure 3 shows the results of gel electrophoresis of six antisense oligonucleotides targeted against the translation initiation site of *bcl-2* mRNA.

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Figures 4(a) and (b) show the degree of DNA fragmentation resulting from oligonucleotide treatment of RS11846 cells. Figure 4(a) shows the effect of oligonucleotides targeted against the translation initiation site. Figure 4(b) shows the effect of oligonucleotides directed against the 5'-cap region of bcl-2 mRNA.

Figure 5 is a graph showing the concentration-dependence of inhibition by an antisense oligonucleotide targeted against the translation initiation site of bcl-2 mRNA.

Figures 6(a) and (b) are graphs showing the results of immunofluorescence analysis of bcl-2 protein levels in oligonucleotide-treated cells.

Figures 7(a-d) are FACS profiles for 697 cells before and after treatment with bcl-2 antisense oligonucleotides.

Figure 8 (a-c) show bcl-2 antisense oligodeoxynucleotides producing sequence-specific reductions in bcl-2 mRNA and bcl-2 protein and producing increased sensitivity of SU-DHL-4 cells to cancer chemotherapeutic drugs.

Figures 9(a) and (b) demonstrate the differential effects of bcl-2 antisense oligomers on chemosensitivity of 32D-bcl-2 and 32D-BHRF-1 cells.

Figures 10 (a-b) show reduction of chemoresistance of RS11846 cells from inducible bcl-2 antisense expression from an expression plasmid.

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Figure 11 shows methylphosphonate/phosphodiester bcl-2 antisense oligomers inducing death of DOHH2 lymphoma cells.

5 Figure 12 shows methylphosphonate (MP)/Phosphodiester (PO) chimeric oligomers inhibiting growth of MCF-7 human breast cancer cells.

Figures 13(a-c)) show optimization of antisense bcl-2 oligomer sequences.

Detailed Description of the Invention

10 According to the invention, anticodc oligomers are provided for inhibiting cancer cell growth, for increasing the sensitivity of cancer cells to cancer chemotherapeutic agents, or for inducing cancer cell death alone or in combination with any one or more cancer
15 chemotherapeutic agents.

Definitions

As used herein, the term "anticode oligomers" means anticodc oligonucleotides and analogs thereof and refers to a range of chemical species that recognize polynucleotide target sequences through hydrogen bonding interactions with the nucleotide bases of the target sequences. The target sequences may be single- or double-stranded RNA or single- or double-stranded DNA.
20
25

The anticodc oligonucleotides and analogs thereof may be RNA or DNA, or analogs of RNA or DNA, commonly referred to as antisense oligomers or antisense oligonucleotides. Such RNA or DNA analogs comprise but are not limited to 2'-O-alkyl sugar modifications, methylphosphonate, phosphorothioate, phosphordithioate, formacetal, 3'-thioformacetal, sulfone, sulfamate, and nitroxide backbone modifications, and analogs wherein the base moieties have been modified. In addition, analogs
30

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of oligomers may be polymers in which the sugar moiety has been modified or replaced by another suitable moiety, resulting in polymers which include, but are not limited to, morpholino analogs and peptide nucleic acid (PNA) analogs (Egholm, et al. Peptide Nucleic Acids (PNA) - Oligonucleotide Analogues with an Achiral Peptide Backbone, (1992)).

Anticode analogs may also be mixtures of any of the oligonucleotide analog types together or in combination with native DNA or RNA. At the same time, the oligonucleotides and analogs thereof may be used alone or in combination with one or more additional oligonucleotides or analogs thereof. The oligonucleotides may be from about 10 to about 1,000 nucleotides long. Although oligonucleotides of 10 to 100 nucleotides are useful in the invention, preferred oligonucleotides range from about 15 to about 24 bases in length.

Anticode oligonucleotides and analogs thereof also comprise conjugates of the oligonucleotides and analogs thereof. (John Goodchild, Conjugates of Oligonucleotides and Modified Oligonucleotides: A Review of Their Synthesis and Properties, Bioconjugate Chemistry, Volume 1 No. 3, May/June (1990)). Such conjugates having properties to improve the uptake, pharmacokinetics, and nuclease resistance of the oligonucleotide, or the ability to enhance cross-linking or cleavage of the target sequence by the oligonucleotide.

As used herein, the term "cell proliferation" refers to cell division rate/cell cycle. The term "growth," as used herein, encompasses both increased cell numbers due to faster cell division and due to slower rates of cell death.

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As used herein, bcl-2 gene expression refers to bcl-2 protein production from the human bcl-2 gene; e.g. reduced bcl-2 gene expression means reduced levels of bcl-2 protein.

5 As used herein, "strategic sites" are defined as any site which when bound by the claimed anticode molecules or analogs thereof results in inhibiting expression of the bcl-2 gene.

10 As used herein, the term "sequence portion" is a portion of the nucleotide sequence of an RNA oligonucleotide. In appropriate contexts, "sequence portion" may refer to a portion of the nucleotide sequence of a DNA segment or DNA oligonucleotide.

15 Uncontrolled cell proliferation is a marker for a cancerous or abnormal cell type. Normal, non-cancerous cells divide regularly, at a frequency characteristic for the particular type of cell. When a cell has been transformed into a cancerous state, the cell divides and proliferates uncontrollably. Inhibition of proliferation 20 modulates the uncontrolled division of the cell. Containment of cell division often correlates with a return to a non-cancerous state.

25 A human gene termed bcl-2 (B cell lymphoma/leukemia-2) is implicated in the etiology of some common lymphoid tumors, Croce et al., "Molecular Basis Of Human B and T Cell Neoplasia," in: Advance in Viral Oncology, 7:35-51, G. Klein (ed.), New York: Raven Press, 1987. High levels of expression of the human bcl-2 gene have been found in all lymphomas with t (14; 30 18) chromosomal translocations including most follicular B cell lymphomas and many large cell non-Hodgkin's lymphomas. High levels of expression of the bcl-2 gene have also been found in certain leukemias that do not

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have a t(14; 18) chromosomal translocation, including most cases of chronic lymphocytic leukemia acute, many lymphocytic leuk mias of the pre-B cell type, neuroblastomas, nasophryngeal carcinomas, and many adenocarcinomas of the prostate, breast, and colon.

(Reed et al., Differential expression of bcl-2 protooncogene in neuroblastoma and other human tumor cell lines of neural origin. Cancer Res. 51:6529 (1991); Yunis et al. Bcl-2 and other genomic alterations in the prognosis of large-cell lymphomas. New England J. Med. 320:1047; Campos et al. High expression of bcl-2 protein in acute myeloid leukemia is associated with poor response to chemotherapy. Blood 81:3091-3096 (1993); McDonnell et al. Expression of the protooncogene bcl-2 and its association with emergence of androgen-independent prostate cancer. Cancer Res. 52:6940-6944 (1992); Lu Q-L, et al. Bcl-2 protooncogene expression in Epstein Barr Virus-Associated Nasopharyngeal Carcinoma, Int. J Cancer 53:29-35 (1993); Bonner et al. bcl-2 protooncogene and the gastrointestinal mucosal epithelial tumor progression model as related to proposed morphologic and molecular sequences, Lab Invest. 68:43A (1993)).

While not limited to the following explanation, the present invention exploits cellular mechanisms concerned with normal cell death. Because most types of cells have a finite life span and are programmed to die, uncontrollable cell accumulation can also result because of a defect in normal cell death mechanisms rather than through an increased rate of cell division. The bcl-2 gene contributes to the pathogenesis of cancer primarily by prolonging cell survival rather than accelerating cell division.

Antisense oligomers suitable for use in the invention include nucleotide oligomers which are two to

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- two hundred nucleotide bases long; more preferably ten to forty bases long; most preferably twenty bases long. The oligonucleotides are preferably selected from those oligonucleotides complementary to strategic sites along the pre-mRNA of *bcl-2*, such as the translation initiation site, donor and splicing sites, or sites for transportation or degradation.

Blocking translation at such strategic sites prevents formation of a functional *bcl-2* gene product. It should be appreciated, however, that any combination or subcombination of anticod oligomers, including oligonucleotides complementary or substantially complementary to the *bcl-2* pre-mRNA or mRNA that inhibit cell proliferation is suitable for use in the invention. For example, oligodeoxynucleotides complementary to sequence portions of contiguous or non-contiguous stretches of the *bcl-2* RNA may inhibit cell proliferation and would thus be suitable for use in the invention.

It should also be appreciated that anticod oligomers suitable for use in the invention may also include oligonucleotides flanking those complementary or substantially complementary to such sequence portions as the strategic or other sites along the *bcl-2* mRNA. The flanking sequence portions are preferably from two to about one hundred bases, more preferably from about five to about twenty bases in length. It is also preferable that the anticod oligomers be complementary to a sequence portion of the pre-mRNA or mRNA that is not commonly found in pre-mRNA or mRNA of other genes to minimize homology of anticod oligomers for pre-mRNA or mRNA coding strands from other genes.

Preferred antisense, or complementary, olig deoxynucleotides are listed in Table 1.

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TABLE I

bcl-2 Oligodeoxynucleotides

	translation initiation antisense (TI-AS)	3'...CCCTCCTACCGCGTGCAGAC...5'
5	<i>bcl-2</i>	5'...CTTTCCCTGGGAAGGATGGCGCACGCTGGGAGA...3'
	splice donor antisense (SD-AS)	31...CCTCCGACCCATCCACGTAG...5'
	<i>bcl-2</i>	5'...ACGGGGTAC...GGAGGCTGGGTAGGTGCATCTGGT...3'
10	splice acceptor antisense (SA-AS)	3'...GTTGACGTCCTACGGAAACA...5'
	<i>bcl-2</i>	5'...CCCCAACTGCAGGATGCCTTGAACTGTACGG...3'

It will be appreciated by those skilled in the art to which this invention pertains, that anticod 15 oligomers having a greater or lesser number of substituent nucleotides, or that extend further along the *bcl-2* mRNA in either the 3' or 5' direction than the preferred embodiments, but which also inhibit cell proliferation are also within the scope of the invention.

It is preferable to use chemically modified 20 derivatives or analogs of anticod oligomers in the performance of the invention rather than "native" or unmodified oligodeoxynucleotides. "Native" oligodeoxynucleotides can be conveniently synthesized 25 with a DNA synthesizer using standard phosphoramidite chemistry. Suitable derivatives, and methods for preparing the derivatives, include phosphorothioate, Stein et al., *Nucl. Acids Res.*, 16:3209-3221 (1988); methylphosphonate, Blake et al., *Biochemistry* 24:6132-6138 (1985) and alphadeoxynucleotides, Morvan et 30 al., *Nucl. Acids Res.* 14:5019-5032 (1986), 2'-O-methyl-ribonucleosides (Monia et al. Evaluation of 2'-modified oligonucleotides containing 2' deoxy gaps as antisense inhibitors of gene expression. *J. Biol. Chem.* 268:14514-14522 (1993)), and covalently-linked derivatives such as

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acridine, Asseline et al., Proc. Natl Acad. Sci. USA 81:3297-3201 (1984); alkylated (e.g., N-2-chlorocethylamine), Knorre et al., Biochemie 67:783-789 (1985) and Vlassov et al., Nucl. Acids Res. 14:4065-4076 (1986); phenazine, Knorre et al., *supra*, and Vlassov et al., *supra*; 5-methyl-N⁴-N⁴-ethanocytosine, Webb et al., Nucl. Acids Res. 14:7661-7674 (1986); Fe-ethylenediamine tetraacetic acid (EDTA) and analogues, Boutorin et al., FEBS Letter's 172:43-46 (1984); 5-glycylamido-1, 10-o-phenanthroline, Chi-Hong et al., Proc. Natl. Acad. Sci. USA 83:7147-7151 (1986); and diethylenetriamine-pentaacetic acid (DTPA) derivatives, Chu et al., Proc. Natl. Acad. Sci. 82:963-967 (1985). All of the above publications are hereby specifically incorporated by reference as if fully set forth herein.

The anticode oligomer of the present invention can also be combined with a pharmaceutically acceptable carrier for administration to a subject or for ex-vivo administration. Examples of suitable pharmaceutical carriers are a variety of cationic lipids, including, but not limited to N-(1-2,3-dioleyloxy)propyl)-n,n,n-trimethylammonium chloride (DOTMA) and dioleoylphophatidylethanolamine (DOPE)]. Liposomes are also suitable carriers for the anticode oligomers of the invention.

The anticode oligomers may be administered to patients by any effective route, including intravenous, intramuscular, intrathecal, intranasal, intraperitoneal, subcutaneous injection, in situ injection and oral administration. Oral administration requires enteric coatings to protect the claimed anticode molecules and analogs thereof from degradation along the gastrointestinal tract. The anticode oligomers may be mixed with an amount of a physiologically acceptable carrier or diluent, such as a saline solution or other

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suitable liquid. The anticode oligomers may also be
combined with liposomes or other carrier means to
protect the anticode molecules or analogs thereof from
degradation until they reach their targets and/or
5 facilitate movement of the anticode molecules or analogs
thereof across tissue barriers.

The anticode oligomers may also be useful for
ex vivo bone marrow purging. Normally, the amounts of
conventional cancer chemotherapeutic agents or drugs and
10 irradiation that a patient can receive are limited by
toxicity to the marrow, i.e., anemia (fatigue, heart
failure), thrombocytopenia (bleeding), neutropenia
(infection). Thus, in order to deliver sufficient
concentrations of drugs and irradiation to totally
15 eradicate the tumor, the physician would simultaneously
destroy the patient's normal bone marrow cells leading to
patient demise. Alternatively, large amounts of bone
marrow can be surgically extracted from the patient and
stored in vitro. While the patient receives aggressive
20 conventional treatment. The patient can then be rescued
by reinfusion of their own bone marrow cells, but only if
that marrow has been "purged" of residual malignant
cells. The claimed anticode oligomers could be used to
remove residual malignant cells from the bone marrow.

25 The anticode oligomers are administered in
amounts effective to inhibit cancer or neoplastic cell
growth. The actual amount of any particular anticode
oligomer administered will depend on factors such as the
30 type of cancer, the toxicity of the anticode oligomer to
other cells of the body, its rate of uptake by cancer
cells, and the weight and age of the individual to whom
the anticode oligomer is administered. Because of
inhibitors present in human serum that may interfere with
35 the action of the anticode oligomer an effective amount
of the anticode oligomer for each individual may vary.

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An effective dosage for the patient can be ascertained by conventional methods such as incrementally increasing the dosage of the anticode oligomer from an amount ineffective to inhibit cell proliferation to an effective amount. It is expected that concentrations presented to cancer cells in the range of about 0.001 micromolar to about 100 micromolar will be effective to inhibit cell proliferation.

The anticode oligomers are administered to the patient for at least a time sufficient to inhibit proliferation of the cancer cells. The anticode oligomers are preferably administered to patients at a frequency sufficient to maintain the level of anticode oligomers at an effective level in or around the cancer cells. To maintain an effective level, it may be necessary to administer the anticode oligomers several times a day, daily or at less frequent intervals. Anticode oligomers are administered until cancer cells can no longer be detected, or have been reduced in number such that further treatment provides no significant reduction in number, or the cells have been reduced to a number manageable by surgery or other treatments. The length of time that the anticode oligomers are administered will depend on factors such as the rate of uptake of the particular oligodeoxynucleotide by cancer cells and time needed for the cells to respond to the oligodeoxynucleotide. *In vitro*, maximal inhibition of neoplastic cell growth by "native," unmodified anticode oligomers occurred two days after initiation of cultures, whereas phosphorothioate oligodeoxynucleotides required 4 to 7 days to achieve maximal inhibition. *In vivo*, the time necessary for maximal inhibition of cell proliferation may be shorter or longer.

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The anticode oligomers of the invention may be administered to patients as a combination of two or more different anticode oligomer oligodeoxynucleotide sequences or as a single type of sequence. For instance, 5 TI-AS and SD-AS could be administered to a patient or TI-AS alone.

It is also believed that the anticode oligomers of the invention may be useful in the treatment of autoimmune diseases. Autoimmune diseases are those 10 diseases in which the body's immune system has malfunctioned in some way. Administration of the anticode oligomers of the invention to a person having an autoimmune disease should inhibit proliferation of bcl-2 overexpressing lymphocytes, which would in turn reduce 15 the symptoms of the autoimmune disease. For use in treating autoimmune diseases, the anticode oligomers would be administered as described herein.

EXAMPLES

General Methods

20 The Examples below use the following protocols:
A. Cells and Cell Cultures. Human leukemic cells lines used for these studies were RS11846 follicular lymphoma cells, 697 pre-B cell acute lymphocytic leukemic cells, and JURAT T cell acute 25 lymphocytic leukemic cells as described in Tsujimoto et al., Proc. Natl. Acad. Sci. USA, 83:5214-5218 (1986) and Weiss et al., Proc. Natl. Acad. Sci. USA, 138:2169-2174 (1987). Human peripheral blood lymphocytes (PBL) were isolated from fresh whole blood as described in Reed et 30 al., J. Immunol., 134:314-319 (1985). All lymphoid cells were cultured at 5×10^5 cells/ml in RPMI medium supplemented with 1 mM glutamine, antibiotics, and either 35 5-10% (v:v) fetal bovine serum (FBS), 5-10% (v:v) calf serum (CS) (both from Hyclone Laboratories), or 1% (v:v) HLI concentrated supplement (Ventrex Laboratories) for

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serum-free cultures. Murine fibroblast cell lines were added at 10^3 cells/cm² in DMEM medium containing glutamine, antibiotics and 5-10% (v:v) FCS. Fibroblast cell lines were NIH 3T3 cells, 3T3-B-alpha-S cells, and 3T3-B-alpha-AS cells. These latter two cell lines are NIH 3T3 cells that express high levels of a human *bcl-2-alpha* cDNA in either the sense or antisense orientation, respectively, by virtue of stable transfection with expression vectors constructs.

B. Measurement of Cellular Growth. Growth of cell lines cultured in the presence or absence of anticod oligomers was measured by two methods: cell counts using a hemocytometer; and DNA synthesis by assaying [³H]-thymidine incorporation essentially as described in Reed et al., *J. Immunol.*, 134:314-319 (1985). Briefly, cells were cultured in 96-well flat-bottomed microtiter plates (Falcon) at 0.2 ml/well. At appropriate times, cells were resuspended, 25 μ l removed from cultures for cell counting, and this volume replaced with 25 μ l of 20 UCi/ML [³H]-thymidine (specific activity 6.7 Ci/mmol) (New England Nuclear). Microtiter cultures were then returned to 37°C and 95% air: 5% CO₂ atmosphere for 8 hours before lysing cells an glass filters and determining relative levels of [³H]-thymidine incorporation into DNA by scintillation counting. Cell counts were performed in the presence of trypan blue dye to determine the concentration of viable cells in duplicate microcultures.

MTT [3-(4, 5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide)] dye reduction assays were performed by the method of Tada, et al. *J. Immunol Methods* 93, 157 (1986), and confirmed to be within the linear range of the assay under the conditions described here. The number of viable cells per well was extrapolated from standard curves that were included with each assay and

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that consisted of serial two-fold dilutions of exponentially growing SU-DHL-4 cells in HL-1 medium, beginning with 10^6 cells/ml (0.2ml/well). Samples were assayed in triplicate and the OD_{600nm} for a media/reagent blank was subtracted from all values prior to calculations.

C. RNA Blot Analysis. Total cellular RNA was isolated by a quanidinium isothiocyanate/phenol procedure as described in Chomczynski et al., *Analyt. Biochem.*, 162:156-159 (1987). The polyadenylated fraction was purified by oligodeoxythymidine-cellulose chromatography as described in Aviv et al., *Proc. Natl. Acad. Sci. USA*, 69:1408-1412 (1972). Approximately 5 µg aliquots of mRNA were size-fractionated in 0.8% agarose/6% formaldehyde gels and transferred to nylon membranes. Blots were prehybridized, hybridized, and washed exactly as described in Reed et al., *Mol. Cell Biol.*, 5:3361-3366 (1985), using either a ³²P-cDNA for human bcl-2, as described in Tsujimoto et al., *Proc. Natl. Acad. Sci. USA*, 83:5214-5218 (1986), or a murine bcl-2 probe, pMBCL5.4 as described in Negrini et al., *Cell*, 49:455-463 (1987). Blots were exposed to Kodak XAR film with intensifying screens at -70°C for 1-10 days. Eluting ³²P-bcl-2 probes from membranes and rehybridizing with a ³²P probe for mouse beta-2-microglobulin verified nearly equivalent amounts of mRNA for all samples on blots.

EXAMPLE 1

Preparation of Anticode oligomers

Normal and phosphorothioate

oligodeoxynucleotides were synthesized using an Applied Biosystems 380B DNA synthesizer, and purified by HPLC reverse-phase chromatography (PRP-1 column) as described in Stein et al., *Nucl. Acids Res.*, 16:3209-3221 (1988) which is specifically incorporated as if fully set forth herein. In some cases it was necessary to further purify

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oligodeoxynucleotides by C18-Sep-Pak chromatography (Waters Associates, Millipore, Inc.), as described previously in Kern et al., *J. Clin. Invest.*, 81:237-244 (1988), to eliminate nonspecific cytotoxic activity.

5 Oligodeoxynucleotides eluted in 30% acetonitrile were evaporated to dryness, resuspended at 1-2 mM in sterile Dulbecco's phosphate-buffered saline or Hanks' buffered salt solution (both from Gibco), and stored at -80°C in small aliquots.

10 Table 1 shows the oligodeoxynucleotides synthesized and their relation to the sense-strand of the human *bcl-2* gene. Portions of the sequence of the coding strand of the human *bcl-2* gene are shown, including the translation initiation site (top), splice donor site
15 (middle), splice acceptor region (bottom), and empirically selected sites within the 5' untranslated portion of *bcl-2* pre-mRNA. The ATG initiation codon, GT splice donor, and AG splice acceptor consensus sequences are in boxes.

20 The sequences of the oligodeoxynucleotides synthesized for these investigations are presented, and their relation to human *bcl-2* mRNA is indicated. The TI-AS oligodeoxynucleotide is antisense at the translation initiation site and TI-S is its complementary
25 sense version. SD-AS and SD-S are oligodeoxynucleotides having antisense and sense orientations, respectively, relative to the splice donor region.

The oligodeoxynucleotide TI-AS straddles the predicted translation-initiation site of *bcl-2* mRNAs and
30 is complementary (antisense) to this region. As a control, the sense version of this 20 bp oligodeoxynucleotide, TI-S, was also synthesized.

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In an effort, to specifically block splicing of *bcl-2* mRNAs, a 20 bp antisense oligodeoxynucleotide, SD-AS, was synthesized that overlaps the splice donor site in *bcl-2* primary transcripts. In addition, a complementary sense oligodeoxynucleotide, SD-S, was prepared as depicted in Table 1. The human *bcl-2* gene gives rise to several transcripts through alternative splice site selections, see Tsujimoto et al., Proc. Natl. Acad. Sci. USA, 83:5214-5218 (1986). The preponderance of these transcripts depend upon splicing and encode a 26 kDa protein, *bcl-2-alpha*. One minor transcript, however, does not undergo a splice and consequently encodes a 22 kDa protein *bcl-2-beta*. The SD-AS oligodeoxynucleotide can thus potentially block maturation of most but not all *bcl-2* transcripts.

EXAMPLE 2

Treatment of Serum for In Vitro Investigations
of Antisense Normal Oligodeoxynucleotides

Because normal oligodeoxynucleotides are sensitive to degradation by nucleases present in serum, the efficacy of the TI-AS oligodeoxynucleotide in fetal bovine serum (FBS) heated for 30 minutes at 56°C (the usual procedure for inactivating serum complement) was contrasted with the efficacy of TI-AS in FBS heated for 1 hour at 68°C, a temperature sufficient for irreversible inactivation of many nucleases. The RS11846 follicular lymphoma cell line was used. RS11846 cells contain a t (14; 18) chromosomal translocation that deregulates *bcl-2* expression, resulting in the accumulation of high levels of *bcl-2* mRNAs, Tsujimoto et al., Proc. Natl. Acad. Sci. USA, 83:5214-5218 (1986).

RS11846 follicular lymphoma cells were cultured in medium containing 5% (vol:vol) fetal bovine serum (FBS) that had been heated at 56°C for 0.5 hours or at 68°C for 1 hour. TI-AS normal oligodeoxynucleotide was

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added at the initiation of cultur , and the density of viable cells determined two days later.

5 The TI-AS normal oligodeoxynucleotide was more effective in 68 C-treated serum at suppressing the growth in culture of these lymphoma cells. In all subsequent experiments, sera heated at 68°C for 1 hour prior to use were used in cultures. This treatment did not impair the growth-supporting capacity of the sera.

EXAMPLE 3

10 Specific Inhibition of Lymphoid Cell Growth by Antisense Normal Oligodeoxynucleotides

15 Antisense normal oligodeoxynucleotides directed against the translation initiation site (TI-AS) and the splice donor site (SD-AS) of *bcl-2* transcripts were tested for their ability to suppress the proliferation of 15 normal and neoplastic lymphoid cells.

20 RS11846 follicular lymphoma cells, JUKRAT T cell leukemia cells, and freshly isolated peripheral blood lymphocytes were cultured in medium containing 10% (vol:vol) FBS that had been heated at 68°C for one hour. 25 various concentrations of normal oligodeoxynucleotides were added at the initiation of culture, including: TI-AS, TI-S, SD-AS, and SD-S. Relative DNA synthesis was measured in cultures after 2-3 days by [³H]-thymidine incorporation. Data were calculated as a percentage of control cultures containing volumes of PBS or HBSS equivalent to oligodeoxynucleotide-treated cultures, and represent the mean (\pm standard deviation) of duplicate cultures.

30 Similar data were obtained by measuring cell counts, excluding cold thymidine inhibition as an explanation for the suppression of DNA synthesis observed in cultures treated with antisense oligodeoxynucleotides.

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As shown in Figures 1(a), (b), (c) and (d) both the TI-AS and SD-AS oligodeoxynucleotides inhibited the growth of RS11846 cells in a concentration-dependent manner. The SD-AS oligonucleotide was less effective in 5 inhibiting cell growth than the TI-AS oligodeoxynucleotide. In contrast to these antisense oligodeoxynucleotides, sense oligodeoxynucleotides (TI-S and SD-S) were not inhibitory even at concentrations of up to 250 µG/ml. Moreover, non-sense 10 oligodeoxynucleotides (i.e., those having the same base composition as the antisense oligodeoxynucleotides but with scrambled sequences) also failed to suppress the proliferation of RS11846 cells. The data thus indicate that antisense oligodeoxynucleotides can specifically 15 block the proliferation of these tumor cells. Several other leukemic cell lines that express the *bcl-2* gene were also tested for inhibition of their proliferation by TI-AS and SD-AS oligonucleotides. As with the JURKAT T cell acute lymphocytic leukemic cells, in every case a 20 specific and concentration-dependent decrease in the growth of these human leukemic cells in cultures containing antisense oligodeoxynucleotides was observed.

It has been demonstrated that *bcl-2* expression is transiently induced in normal human peripheral blood 25 lymphocytes (PBL) when these cells are stimulated to proliferate, suggesting that this gene may play a role in the regulation of normal lymphocyte growth, Reed et al., *Science* 236:1295-1297 (1987). The capacity of antisense oligodeoxynucleotides to impair the growth of PBL cultured with a monoclonal antibody, OKT3 (Van den Elsen 30 et al., *Nature* 312:413-418 (1984)), that stimulates their proliferation was therefore tested. PBL were stimulated with 50 µl of purified OKT3 monoclonal antibody. As shown in Figures 1 (a), (b), (c) and (d) the TI-AS 35 oligodeoxynucleotide specifically suppressed the proliferation of PBL in a concentration-dependent manner.

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These antisense normal oligodeoxynucleotides thus suppressed the growth in culture of leukemic cells that constitutively express the *bcl-2* gene and of normal lymphocytes where *bcl-2* expression is inducible.

5

EXAMPLE 4

Time-Course of Inhibition by
Antisense Normal Oligodeoxynucleotides

The kinetics of inhibition by antisense oligodeoxynucleotides was examined in cultures of RS11846 follicular lymphoma cells and of 697 pre-B cell acute lymphocytic leukemia cells. Both of these neoplastic B cell lines transcribe and accumulate *bcl-2* mRNAs at high levels, Tsujimoto et al., Proc. Natl. Acad. Sci. USA, 83:5214-5218 (1986).

15

RS11846 follicular lymphoma and 697 pre-B cell leukemia cells were cultured in medium containing 10% (vol:vol) 68°C-treated FBS and normal oligodeoxynucleotides. Cells were cultured with 50 µg/ml TI-AS, 100 µg/ml SD-AS, 50 µg/ml TI-S (RS11846 cells) or 20 100 µg/ml SO-S (697 cells), or PBS as a control. DNA synthesis (kcpm/10⁵ viable cells) and cell densities (10⁵ viable cells/ml) were measured at various times after initiation of cultures.

25

Antisense normal oligodeoxynucleotides markedly inhibited DNA synthesis measured in cultures of these cells within 24 hours. Diminished cell densities were readily apparent in these cultures within 2 days. Antisense normal oligodeoxynucleotides thus rapidly inhibited the *in vitro* growth of leukemic cells. The 30 action of antisense oligodeoxynucleotides was specific, since sense oligodeoxynucleotides did not impair proliferation in these cultures. Though cell viabilities often declined during the later days of culture no increase in cell death was seen during the first 1-2 days

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of culture with antisense oligodeoxynucleotides,
suggesting a non-cytotoxic mechanism.

EXAMPLE 5

Comparision of Different Serum Preparations

5 Inhibition of proliferation of leukemic cells
with antisense oligodeoxynucleotides can vary greatly
depending on the lot of serum used in cultures.

10 To determine the effects of serum of inhibition
of proliferation, relative levels of DNA synthesis were
measured in cultures of 697 pre-B cell leukemia cells
2 days after addition of 200 µM TI-AS normal
oligodeoxynucleotide. Cells were cultured in medium
supplemented with 1% (vol:vol) HL1-concentrate
(serum-free condition), 5% (vol:vol) of two different
15 lots of calf serum (CS1 and CS2), or 5% (vol:vol) of two
different lots of fetal bovine serum (FBS1 and FBS2).
All sera were heated at 68°C for 1 hour prior to use in
cultures.

20 The normal TI-AS oligodeoxynucleotide markedly
inhibited DNA synthesis (92%) and cellular proliferation
in serum-free cultures (HL1) of 697 cells. This
antisense oligodeoxynucleotide was equally effective
(94%) in cultures containing 5% (v:v) of one of the lots
of fetal bovine serum (FBS2). In contrast, inhibition
25 was significantly reduced in cultures containing other
serum preparations (CS1, CS2, FBS1). It has been
generally observed that antisense normal
oligodeoxynucleotides are less effective in cultures
supplemented with calf serum (CS) than in those
30 containing fetal bovine serum (FBS).

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EXAMPLE 6

Concentration Dependence of Inhibition by Antisense
Normal Oligodeoxynucleotides in Serum-Free Cultures

697 pre-B cell leukemia cells were cultured in
5 medium with either 1% (vol:vol) HL1-concentrate
(serum-free conditions or 5% (vol:vol) 68°C-treated
FBS2). Relative levels of DNA synthesis and cellular
densities measured after 2 days in cultures containing
various concentrations of normal TI-AS
10 oligodeoxynucleotide.

The TI-AS oligodeoxynucleotide was inhibitory
at lower concentrations when used in serum-free cultures.
At 100 μ M, for instance, no inhibition of cellular
proliferation was seen in FBS2-containing cultures,
15 whereas cell counts were reduced by approximately 75% in
serum-free cultures. At higher concentrations of
antisense oligodeoxynucleotides (200-250 μ M), however,
inhibition of 697 cellular proliferation was comparable
in both types of cultures. The increased efficacy of
20 normal oligodeoxynucleotides in serum-free cultures was
specific, since the sense oligonucleotide (TI-S) was not
inhibitory at the same concentrations.

EXAMPLE 7

Antisense Phosphorothioate Oligodeoxynucleotides:

Time Course of Inhibition

To contrast the efficacy of phosphorothioate
oligodeoxynucleotides with that of normal
oligodeoxynucleotides with regard to inhibition of human
leukemic cell growth, phosphorothioate
30 oligodeoxynucleotides were cultured with 697 pre-B cell
leukemia cells and the effects on inhibition were
measured. 697 pre-B cell leukemia cells were cultured in
serum-free medium for various times before measuring DNA
synthesis (kcpm) and cell densities (10^6 cells/ml). Cells
35 were seeded at an initial density of either 0.2×10^5

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- cells/ml or 0.5×10^5 cells/ml. Culture conditions were
- 25 μ M TI-AS phosphorothioate , 25 μ M TI-S phosphorothioate, and control cultures treated with HBSS.

To avoid experimental variation due to
5 differences among lots of sera, 697 leukemic cells were
cultured in serum-free conditions. When cultured at an
initial seeding density of 0.5×10^6 cells/ml, 697 cells
achieved maximal DNA synthesis and cellular densities at
4-5 days. Addition of 25 μ M sense phosphorothioate
10 oligodeoxynucleotide (TI-S) at the initiation of these
cultures had little effect on 697 cell growth. In
replicate cultures containing 25 μ M antisense
phosphorothioate (TI-AS), however, some diminution in DNA
synthesis was evident within 2 days and was maximal at
15 4-5 days. Maximal inhibition of 697 cell growth, as
determined by cell counts, was seen at 6 days after
initiation of cultures.

When 697 cells were initially seeded at 0.2×10^6
cells/ml, the antisense phosphorothioate
20 oligodeoxynucleotide, TI-AS, resulted in only slight
inhibition at 2 days, attaining maximal suppression of
DNA synthesis in these cultures at day 7. As with normal
oligodeoxynucleotides, this inhibition by
phosphorothioate oligodeoxynucleotides appeared to be
25 mediated through non-cytotoxic mechanisms, since cellular
viabilities did not decline until late in the course of
culture. Compared with normal antisense
oligodeoxynucleotides, therefore, phosphorothioate
oligodeoxynucleotides had a slower onset of action.

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EXAMPLE 8

Concentration Dependence of Inhibition by Antisense bcl-2 Phosphorothioate Oligodeoxynucleotides

5 The concentration dependence of inhibition by phosphorothioate and normal TI-AS oligodeoxynucleotides in cultures of 697 cells in serum-free medium was compared as follows.

10 697 cells were cultured in serum-free medium for either 3 days (normal oligodeoxynucleotides) or 4 days (phosphorothioate oligodeoxynucleotides) prior to measuring cell densities and levels of DNA synthesis. Oligodeoxynucleotide additions to cultures included TI-AS phosphorothioate, TI-S phosphorothioate, TI-AS normal, and TI-S normal.

15 As shown in Figures 2 (a) and (b), TI-AS phosphorothioate oligodeoxynucleotides markedly inhibited the proliferation of 697 cells at 25-50 μ M. In contrast, normal TI-AS oligodeoxynucleotides required concentrations 5- to 10-fold higher (approximately 250 μ M) to cause a comparable suppression of 697 cellular proliferation. Suppression by the antisense phosphorothioate oligodeoxynucleotide TI-AS was specific over this concentration range, since its complementary sense oligodeoxynucleotide (TI-S) produced little inhibition of 697 cell growth in replicate cultures (see Figures 2(a) and (b)).

EXAMPLE 9

Influence of Serum Preparation on Inhibition by Antisense Phosphorothioate Oligodeoxynucleotides

30 To further define the effects of serum preparation on the inhibitory activity of phosphorothioate oligodeoxynucleotides, FBS that had been heated to 56°C for 30 minutes, 68°C for 1 hour, or not

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heated prior to addition to cultures was added to cultures of RS11846 lymphoma cells.

RS11846 cells were cultured in medium containing 1% (vol:vol) HL1-concentrate or 5% (vol:vol) FBS that had been heated at 56°C for 0.5 hour, 68°C for 1 hour, or that had not been heated. Cell counts were calculated as a percentage relative to control cultures treated with equivalent concentrations of TI-S phosphorothioate oligodeoxynucleotide, and represent the mean percentage (standard deviation was less than 10% for all values) for duplicate cultures counted on days 4 and 5.

The TI-AS phosphorothioate oligodeoxynucleotide completely inhibited the growth of RS11846 cells at 25 µM, with an estimated half-maximal inhibitory concentration of approximately 11 µM. In contrast, this phosphorothioate oligodeoxynucleotide was considerably less effective in cultures containing 5% (v:v) FBS. Furthermore, heating FBS prior to adding it to cultures did not significantly improve the ability of the TI-AS phosphorothioate oligodeoxynucleotide to suppress the growth of RS11846 lymphoma cells. At an oligodeoxynucleotide concentration of 50 µM, inhibition of proliferation of RS11846 cells never exceeded 48% serum-containing cultures, regardless of the heating procedure used.

EXAMPLE 10

Influence of Dialysis of Serum on Inhibition by Normal and Phosphorothioate Antisense Oligodeoxynucleotides

To further characterize the nature of the interfering substances in serum, experiments were performed wherein 68°C-heated serum was extensively dialyzed (molecular weight cutoff = 3500) prior to being added to cultures of 697 leukemic cells. Experiments were conducted with 12.5 µM TI-AS phosphorothioate

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oligode oxyribonucleotide and 200 μ M of the normal oxygen-based TI-AS oligodeoxyribonucleotide.

697 cells were cultured in medium containing 1% (vol:vol) HL1-concentrate (A) or 5% (vol:vol) of three different lots of 68°C-treated FBS (B,C,D). Each serum preparation was contrasted before (ND) and after (D) extensive dialysis. TI-AS (+) and TI-S (-) oligodeoxyribonucleotides were added to replicate cultures at 200 μ M for normal oxygen-based oligodeoxyribonucleotides (OXY) and at 12.5 μ M for phosphorothioate oligodeoxyribonucleotides (PT). Relative levels of DNA synthesis (kcpm) were measured after 2 or 4 days of culture for normal and phosphorothioate oligodeoxyribonucleotides, respectively.

For the three different lots of FBS tested, two exhibited little change after dialysis in cultures containing either normal or phosphorothioate oligodeoxyribonucleotides. One lot of FBS, however, appeared to interfere less with the inhibitory activities of these antisense oligodeoxyribonucleotides after dialysis.

EXAMPLE 11

Experiments with Stably Transfected NIH 3T3 Cells

Though the antisense oligodeoxyribonucleotides described herein were designed to block bcl-2 mRNA translation (TI-AS) and splicing (SD-AS), the molecular mechanisms of their actions are not yet known. To determine the effect of formation of oligodeoxyribonucleotide-RNA hybrids within cells upon inhibition of cellular growth, irrespective of the nucleotide sequence, cells transformed to express human bcl-2 cDNA transcripts were cultured with normal oligodeoxyribonucleotides.

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- 200 μ M of normal TI-AS and TI-S oligodeoxynucleotides were added to cultures of typical NIH 3T3 cells and to cultures of these cells that had been stably transfected with expression constructs that produce high levels of human *bcl-2* cDNA transcripts for either the usual sense (3T3-alpha-S cells) or the antisense (3T3-alpha-AS cells) strand.

For RNA blot analyses, polyadenylated mRNA was purified from normal NIH 3T3 cells and from cells stably transfected with expression constructs that produce either sense (3T3-alpha-S) or antisense (3T3-alpha-AS) recombinant *bcl-2*-alpha mRNAs, according to the method of 13. Approximately 5 μ g of mRNA was subjected to RNA blot analysis, essentially as described in (16), using either 15 32 P-labeled hybridization probes derived from human or murine *bcl-2* sequences.

An autoradiogram resulting from a one-day exposure of a blot containing RNAs from normal 3T3 cells, 3T3-alpha-AS cells, and 3T3-alpha-S cells showed high relative levels of recombinant 2.4 and 1.4 kbp *bcl-2* transcripts produced from the *bcl-2* expression constructs that were transfected into 3T3-alpha-AS and 3T3-alpha-S cells.

A 10-day exposure of a blot containing RNA from normal 3T3 cells that were either proliferating or quiescent at the time of harvesting RNA showed low but detectable levels of normal 7.5 and 2.4 kbp murine *bcl-2* transcripts present in proliferating 3T3 cells.

TI-AS oligodeoxynucleotide specifically suppressed DNA synthesis and cellular replication in cultures of normal NIH 3T3 cells, consistent with findings by others that fibroblasts do contain *bcl-2* transcripts, albeit at low levels. The TI-AS

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- oligodeoxynucleotide disclosed herein is complementary to the mouse *bcl-2* sequence in 18 of its 20 bases (17), accounting for its ability to suppress the growth of murine NIH 3T3 cells.

5 NIH 3T3 cells, 3T3-alpha-AS cells, and
3T3-alpha-S cells were cultured in medium containing 5%
(vol:vol) 68°C-treated serum and either HBSS, 200 µM TI-S
normal oligodeoxynucleotide, or 200 µM TI-AS normal
oligodeoxynucleotide. Relative levels of DNA synthesis
10 (kcpm) were measured in cultures after 3 days and reflect
a 16 hour incubation with 0.5 µci/well of [³H]-thymidine.
Cell densities, estimated by phase microscopy, were
consistent with the measured DNA synthesis in cultures.
The percentage of inhibition of DNA synthesis in cultures
15 containing TI-AS oligodeoxynucleotides was calculated
relative to control cultures containing HBSS.

As with normal NIH 3T3 cells, culturing
3T3-alpha-S cells (producing human *bcl-2*-alpha sense
transcripts) with TI-AS and TI-S oligodeoxynucleotides
20 demonstrated specific suppression, since the sense
oligodeoxynucleotide TI-S was not inhibitory. The level
of inhibition of cellular proliferation by the antisense
oligodeoxynucleotide, however, was not as great in
3T3-alpha-S cells, as might be expected, since these
25 cells contain more *bcl-2* mRNA.

Adding TI-S oligodeoxynucleotide to cultures of
3T3-alpha-AS cells (produce antisense *bcl-2* transcripts)
ruled out inhibition of cellular growth through a
nonspecific mechanism involving oligodeoxynucleotide--RNA
30 hybrid formation. The TI-S oligodeoxynucleotide caused
little suppression of 3T3-alpha-AS cell proliferation,
whereas the TI-AS oligodeoxynucleotide was markedly
inhibitory in these cells. Similar data were obtained

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with TI-AS and TI-S phosphorothioate oligodeoxynucleotides.

EXAMPLE 12

5 Measurements of DNA Fragmentation as an Indicator of
bcl-2 Antisense Oligodeoxynucleotide-Mediated Programmed
Cell Death in Human Lymphoma Cells

Oligonucleotides having the sequences shown in Table 2 were tested for the ability to induce programmed cell death (DNA fragmentation) in the human 10 t(14:18)-containing human lymphoma cell line RS11846. The oligonucleotides were all phosphodiesters, and were targeted against the translation initiation site or the 5'-cap region of bcl-2 pre-mRNAs. Control 15 oligodeoxynucleotides included a bcl-2 sense version (TI-S) of TI-AS (having SEQ ID NO: 7) and a scrambled version of TI-AS that has the same base composition, but with jumbled nucleotide order.

TABLE 2

	<u>SEQUENCE</u>	<u>SEQ ID NO:</u>
20	CGCGTGCAGAC CCTCTTG	8
	TACCGCGTGC GACCCTC	9
	CCTTCCTAACG GCGTGCG	11
	GACCCTTCCT ACCGCGT	12
	GGAGACCCCTT CCTACCG	13
25	GCGGCCGCAG CGCGG	14
	CGGGCGGGCG ACAGA	15
	CGGGAGCGCG GCAGGC	16

RS11846 cells were adapted to grow in HL1 media with 1% FCS and their DNA metabolically labeled by 30 addition of ¹²⁵I-deoxyuridine to cultures for three hours. Labeled cells were then washed thoroughly and cultured for two days in the presence of various oligonucleotides at 50 AM. Cells were then recovered from 200 μ L cultures by centrifugation, and lysed in a hypotonic buffer

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containing 10 mM EDTA and 1% Triton X100. After centrifugation at 16,000 xg to pellet unfragmented genomic DNA, the supernatant fraction containing fragmented DNA was extracted with phenol/chloroform and 5 ethanol precipitated. This DNA was then subjected to gel electrophoresis in 1.5% agarose gel and transferred to nylon membranes for autoradiography.

The results of two experiments are shown in Figures 3 and 4(a-b). The six bcl-2 antisense 10 oligonucleotides targeted in the vicinity of the ATG site of translation initiation in bcl-2 mRNAs were tested. "C-Oligo-2" refers to an oligonucleotide with 4 purposeful mismatches. "U" indicates untreated control cells. Figures 4 (a-b) show the results for the 15 oligonucleotides shown in Figure 3. "Sc20" refers to a 20mer with the same base composition as TI-AS, but with scrambled sequence. Figure 4(b) shows the results for three oligonucleotides targeted against the 5'-cap of bcl-2 mRNAs. The numbers 20 refer to the distance of these oligomers from the ATG-translation initiation site.

The presence of a ladder of DNA fragments (unit size of approximately 200 bp) is indicative of programmed cell death. At 50 µM, TI-AS caused little DNA 25 fragmentation, whereas the oligonucleotides having SEQ ID NO: 9 and SEQ ID NO: 10, and one of the 5'-cap oligonucleotides (SEQ ID NO: 14) led to pronounced DNA fragmentation.

EXAMPLE 13

30 Concentration-Dependence of Inhibition by Antisense Phosphodiester Oligodeoxynucleotides in Serum-Free Cultures

697 pre-B cell leukemia cells were cultured in medium with either 1% (vol:vol) HL-1 concentrate

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(serum-free conditions [o] or 3% (vol:vol) 68°C-treated serum (FBS2) [__], see Figure 5. Shown are cellular densities measured after 2 days in cultures containing various concentrations of phosphodiester TI-AS oligodeoxynucleotide. Data are shown as percentages relative to control cultures treated with a sense oligonucleotide, and reflect the mean ± standard deviation for duplicate samples.

EXAMPLE 14

10 Immunofluorescence Analysis of bcl-2 Protein Levels in Oligodeoxynucleotide-Treated 697 Cells

For studies with oligodeoxynucleotides, 0.25×10^4 (for phosphorothioate) or 0.5×10^5 (for normal oligodeoxynucleotides), 697 cells were cultured in 1 ml of HL-1 serum-free medium in 24 well culture dishes (Linbro. Flow Lab, Inc.). After 2 days (for normal) or 4 days (for phosphorothioates), cells were recovered from cultures, washed once in [PBS, pH 7.4 (Gibco) - 0.1% bovine serum albumin - 0.1% sodium azide], and fixed for 5-10 minutes on ice in 1% paraformaldehyde/PBS solution. The cells were then washed once in PBS and incubated in 1 ml of absolute methanol at 20°C for 10 minutes. After washing once in PBS-A, cells were then resuspended in PBS containing 0.05% Triton-X100 for 3 minutes on ice, washed in PBS-A and preblocked for 30 minutes at 4°C in PBS with 10% (v/v) heat-inactivated goat serum.

For addition of the first antibody, preblocked cells were resuspended in 100 µl of PBS-G (PBS-1% goat serum-0.1% sodium azide) prior to aliquoting 50 µl into 30 separate tubes that contained 1 µl of either BCL2 antibody (Halder et al., Nature (London), 342:195-197 (1989)) or affinity-purified normal rabbit control IgG (Cappel 6012-0080) and incubated for 1 hour on ice. The BCL2 antibody used for these studies was prepared in 35 rabbits using a synthetic peptide corresponding to amino

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acids (98-114) of the BCL2 protein and was affinity--
purified by protein-A-Sepharose chromatography and used
at approximately 1 mg/ml. Cells were then washed in
5 PBS-A and incubated in 0.5-1.0 ml PBS-A for 15-20 minutes
on ice to allow diffusion of nonspecific cell-associated
antibody prior to resuspending cells in 100 µl of PBS-G
containing 5 µg of biotinylated scat anti-rabbit IgG
(BAI000; Vector Labs) for 30 minutes. After washing once
and incubating for 15 minutes in PBS-A, cells were
10 finally resuspended in 100 µl of PBS-A containing 2 µg of
FITC-conjugated avidin (Vector Labs A2011) for 20 minutes
and washed three times in PBS-A prior to analysis with an
Ortho cytofluorograph 50-H connected to an Ortho 2150
data-handling system. The specificity of method for
15 detecting BCL2 protein was confirmed by
immunofluorescence microscopy (showing cytosolic stain
peptide competition, and studies of cell lines that
expressed various levels of BCL2 mRNA and proteins
through gene transfer manipulations.

20 For measurements of surface HLA-DR antigen
expression, an indirect immunofluorescence assay method
was used (Reed et al., *J. Immunol.* 134:1631-1639 (1985))
involving incubation of viable cells with a murine
anti-HLA-DR monoclonal antibody (IgG2a) (Becton-Dickinson
25 7360) or a negative control antibody, R3-367 (IgG2a),
followed by FITC-conjugated scat anti-mouse IgG (Cappel
1711-0081). Cells were fixed in 1% paraformaldehyde/PBS
prior to FACS analysis.

697 cells were cultured for 2 days (PO) or
30 4 days (PS) with various oligonucleotides. In
Figures 6(a-b), the black columns show the results with a
sense oligonucleotide, and the hatched columns with an
antisense oligonucleotide TI-AS. Cells were labeled with
anti-bcl-2 antiserum and analyzed by FACS. Data are

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expressed as percentages relative to the mean fluorescence obtained with untreated 697 cells.

Figures 7(a-d) show typical FACS results obtained for 697 cells before and after treatment with 5 $100 \mu\text{M}$ PO *bcl-2* antisense oligonucleotides. A: untreated 697 cells labeled with either anti-*bcl-2* antiserum (hatched area) or normal rabbit serum control (white area); B: untreated 697 cells labeled with either anti-HLA-DR antibody (hatched area) or a negative control antibody (white area); C: 697 cells cultured for 2 days 10 with either normal *bcl-2* TI-AS (white area) or TI-AS (hatched area) oligodeoxynucleotides and labeled with anti-*bcl-2* antibody; D: 697 cells cultured with TI-AS and TI-S oligodeoxynucleotides (as in C), but labeled 15 with anti-HLA-DR antibody.

As shown in Figures 6 (a) and (b) , PO and PS *bcl-2* antisense oligonucleotides produced specific concentration-dependent reductions in the levels of *bcl-2* proteins, without altering the levels of expression of 20 HLA-DR (Figures 7(a-d)) and other control antigens. At $150 \mu\text{M}$, for example, PO antisense oligodeoxynucleotide caused an approximately 75-95% reduction in *bcl-2* fluorescence, whereas the control sense oligodeoxynucleotide diminished *bcl-2* protein levels by 25 only 10-20% (Figure 6(a)). Similarly, cultured 697 cells for 4 days with the PS antisense oligodeoxynucleotide at $25 \mu\text{M}$ resulted in approximately 70% reduction in *bcl-2* fluorescence. In comparison, the sense PS oligodeoxynucleotide TI-AS inhibited *bcl-2* protein levels 30 by only approximately 15%, as measured by this assay (Figure 6(b)).

SIGNIFICANCE

In phosphorothioate oligodeoxynucleotides, one of the non-bridging oxygen atoms in each internucleotide

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phosphate linkage is resolaced by a sulfur atom. This modification renders phosphorothioate oligodeoxynucleotides extremely resistant to cleavage by nucleases, Stein et al., *Nucl. Acids Res.*, 16:3209-3221 (1988). Despite the substitution of a sulfur atom for an oxygen, phosphorothioate oligodeoxynucleotides retain good solubility in aqueous solutions; hybridize well, though with some decrease in the melting temperature of RNA-oligodeoxynucleotides duplexes; and are synthesized conveniently by the widely employed method of automated oligodeoxynucleotides synthesis with phosphoroamidites.

Antisense *bcl-2* phosphorothioate

oligodeoxynucleotides have been found to be more potent inhibitors of leukemic cell growth than their normal oxygen-based counterparts. When tested under serum-free conditions, these oligodeoxynucleotides reduced cellular proliferation by half at concentrations of approximately 15-23 μ M, whereas the normal oligodeoxynucleotide achieved 50% inhibition at 125-250 μ M. This finding may be explained by the reduced sensitivity of phosphorothioate oligodeoxynucleotides to cellular nucleases, or may be attributable to other mechanisms. For example, mRNAs hybridized with phosphorothioate oligodeoxynucleotides may experience enhanced degradation through a mechanism involving an RNase H-like activity.

Despite their increased inhibitory activity, phosphorothioate antisense oligodeoxynucleotides retained sequence-specificity. At the concentrations tested (less than 25 μ M), sense versions of these oligodeoxynucleotides had little effect on leukemic cell growth. Both normal and phosphorothioate antisense oligodeoxynucleotides appeared to initially suppress the proliferation of leukemic cells through non-cytotoxic mechanisms. During the first few days of culture, cellular replication was inhibited without a concomitant

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rise in cell death. Later in these cultures (days 4-5 for normal oligodeoxynucleotides, days 6-8 for phosphorothioates), however, cellular viabilities declined.

5 Comparing the kinetics of inhibition by normal and phosphorothioate oligodeoxynucleotides revealed that the latter compounds have a slower onset of action. Maximal inhibition of leukemic cell proliferation by normal antisense oligodeoxynucleotides occurred two days
10 after initiation of cultures, whereas phosphorothioate oligodeoxynucleotides required 4 to 7 days to achieve maximal inhibition.

15 The usefulness of anticod oligomers in inhibiting human lymphoma/leukemia cells and other types of cancer cells that express the *bcl-2* gene has been shown by the examples herein. Anti-sense oligodeoxynucleotides complementary to at least an effective portion of the mRNA of the human *bcl-2* gene has been found to inhibit growth of RS11846 human follicular
20 lymphoma cells t (14;18) chromosomal translocation and high *bcl-2* expression), 697 human pre B cell leukemia cells (high *bcl-2* expression), JURKAT human acute lymphocytic leukemia cells (medium *bcl-2* expression), normal human lymphocytes (medium *bcl-2* expression) and
25 murine fibroblasts (low *bcl-2* expression). Although *bcl-2* antisense reagents can suppress the growth of many types of cells, the t(14;18) lymphoma and leukemia cells seem to be the sensitive, allowing for specific inhibition of malignant cells.

30 As demonstrated in the following Examples, a variety of DNA analogs can be employed in the instant invention. For example, phosphorothioates, methylphosphonates, and mixed oligomers containing combinations of phosphodiesters and phosphorothioate or

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methylphosphonate nucleosides. It shoud be underst od
that RNA analogs can also be employed in the invention.

EXAMPLE 15

Methylphosphonate (MP)/Phosphodiester (PO) bcl-2

5 Antisense Oligomers Induce Death of DoHH2 Lymphoma Cells

The purpose of this study was to determine the efficacy of various analogs of the anticodc oligomers for inhibiting lymphoma cell survival.

DoHH2 is a human lymphoma cell line containing
10 a t(14:18)-translocation that activates the bcl-2 gene.
DoHH2 cells were cultured for 3 days without oligomers or
in the presence of various concentrations of antisense
15 (As) and scrambled (Sc) methylphophonate
(MP)/Phosphodiester (PO) oligomers for 3 days. Cell
viability was assessed by trypan blue dye exclusions, and
the data expressed as a percentage relative to DoHH2
cells cultured without oligomers. The MP/PO oligomers
was an 18-mer targeted against the first 6 codons of the
20 bcl-2 open reading frame in which 5 internal linkages
were phosphodiester and the flanking nucleosides were
methylphophonates.

The results indicate that these anticodc
oligomer analogs are potent and specific inhibitors of
lymphoma cell survival.

25

EXAMPLE 16

Methylphosphonate (MP)/Phosphodiester (PO) Chimeric
Oligomers Inhibit Growth of MCF-7
Human Breast Cancer Cells

The purpose of this study was to determine the
efficacy of the claimed anticodc oligomer analogs to
inhibit the survival of solid tumor cells which highly
30 express bcl-2.

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MCF-7 is a human breast adenocarcinoma cell line
• that contains relatively high levels of bcl-2 protein.
The cells were cultured at 4,000 cells per well in 96-well microtiter plates in the presence or absence of
5 MP/PO oligomers. Relative cell numbers per well were then estimated by MTT assay, based on a standard curve prepared using freshly plated, untreated MCF-7 cells. The antisense (As) and scrambled (Sc) MP/PO oligomers were the same as those described in Example 15. Data
10 represent the mean +/- standard deviation for determinations.

The results demonstrate sequence specific inhibition of growth of solid tumor cells by the claimed anticode oligomer analogs.

15

EXAMPLE 17

Optimization of Anticode bcl-2 Oligomer Sequences

The purpose of this study was to determine optimum target sites or sequence portions on mRNA for inhibiting cell survival by contacting the cells with 20 various claimed anticode molecules whose sequences were computer generated.

DoHH2 lymphoma cells were treated with various concentrations of oligomers targeted to different sites on the bcl-2 mRNA. The ATG oligomer targets the 25 translation initiation site, and is complementary to the first 6 codons of the open reading frame. The Dscore 23 and Dscore 72 oligomers target sites in the 5' untranslated region of the mRNA. Sc oligomers represent negative controls having the same length and base 30 composition but in scrambled order. All oligomers were prepared as phosphodiester (PO)/phosphorothioate (PS) chimeras, where only the last (3') two internucleoside linkages were phosphorothioates. Oligomers were added directly to cultures and relative numbers of viable cells

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were estimated by MTT assay 3 days later. Data represent mean +/- standard deviation.

The results indicate that the Dscore 23 oligomer, targeted to the 5' untranslated region, has, compared to the other anticodc oligomers tested in this Example, superior activity for inhibiting cell survival.

EXAMPLE 18

Reversal of Chemoresistance of Tumor Cells by Antisense-Mediated Reduction of bcl-2 Gene Expression

10 The following work was undertaken to determine if anticodc oligomers directed against the expression of the bcl-2 gene would reverse chemoresistance, that is to say, increase the sensitivity to cancer chemotherapeutic agents in cancer tumor cells expressing the bcl-2 gene.

15 High levels of bcl-2 protein appeared to increase the relative resistance of lymphoid cells to killing induced by a wide variety of cancer chemotherapeutic agents including, but not limited to, Ara-C, MTX, vincristine, taxol, cisplatin, adriamycin, etoposide, mitozantron, 2-chlorodeoxyadenosine, dexamethasone (DEX), and alkylating agents. (Miyashita, T. and Reed, J.C., Cancer Res. 52:5407, October 1, 1992). While these drugs have diverse biochemical mechanisms of action, it is believed that all have in common the ability to ultimately trigger cancer cell death by activating endogenous cellular pathways leading to apoptosis (Eastman, A. Cancer Cells 2:275 (1990)). It is understood that the claimed anticodc molecules and analogs thereof as used herein are effective for their intended purposes of enhancing sensitivity to cancer chemotherapeutic drugs including, but not limited to, antimetabolites, alkylating agents, plant alkaloids, and antibiotics.

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Antimetabolites include, but are not limited to, methotrexate, 5-fluoruracil, 6-mercaptopurine, cytosine arabinoside, hydroxyurea, 20chlorodeoxyadenosine.

5 Alkylating agents include, but are not limited to, cyclophosphamide, melphalan, busulfan, cisplatin, paraplatin, chlorambucil, and nitrogen mustards.

Plant alkaloids include, but are not limited to, vincristine, vinblastine, VP-16.

10 Antibiotics include, but are not limited to, doxorubicin (adriamycin), daunorubicin, mitomycin c, bleomycin.

15 Other cancer chemotherapeutic agents include DTIC (decarbazine), mAMSA, hexamethyl melamine, mitroxantrone, taxol, etoposide, dexamethasone.

In the present work, both nuclease resistance phosphorothioates (PS) and phosphodiesters in which only the 3'-most internucleoside bond was a thioate linkage (PO/PS) were employed. The PO/PS oligomers are 20 resistance to 3' exonucleases (the principal nuclease activity of serum) and generally form more stable heteroduplexes with target RNAs.

Cationic lipids were used to improve the uptake and subsequent release of oligomers into effective 25 intracellular compartments, and are exemplary pharmaceutical carriers for the claimed anticode oligomers.

The methods for preparing and purifying the antisense (AS) and scrambled (SC) 18'mer oligonucleotides 30 used for the present work are described above in General

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Methods and in Kitada et al. (Antisense R & D, 3:157 (1993)). Phosphodiester oligonucleotides were synthesized in a 10-15 micromole scale using phosphoroamidate chemistry with oxidation by iodine, and then purified using a C₁₈-reverse phase column. In most cases, oligomers were additionally ethanol-precipitated five times to eliminate any nonspecific cytotoxic activity, and then dried and resuspended in sterile HL-1 medium (Ventrex Labs, Inc; Burlingame, CA) at 1-10 mM. The pH of this solution was adjusted using 1-10 M NaOH until the phenol red indicator dye in the media returned to its original color.

The principal oligomers used were 18-mers, having either the sequence:

- I. TCTCCCAGCGTGCGCCAT (SEQ ID NO. 17), which is antisense to the first six codons of the human bcl-2 open reading frame (SEQ ID NO. 19); or
- II. TGCACTCACGCTCGGCCT (SEQ ID NO. 18), which is a scrambled version used as a control.

Standard transfection methods were used to produce tumor cells expressing either the bcl-2 gene or an antisense oligodeoxynucleotide which binds to bcl-2 mRNA. It is understood that the vector could also encode an antisense oligodeoxynucleotide which binds to bcl-2 pre-mRNA. The particular nucleotide sequence encoding the antisense oligonucleotides of the invention is not critical, except that the sequences are preferably chosen such that they express antisense oligodeoxynucleotides sufficient to reduce bcl-2 gene expression in tumor cells and increase the sensitivity of the tumor cells to cancer chemotherapeutic agents or sufficient to kill tumor cells when they are treated with cancer chemotherapeutic agents. It is only necessary that the antisense oligodeoxynucleotide encoded in vector is expressed under

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conditions sufficient to reduce bcl-2 gene expression in tumor cells. The methods used for preparing vectors, and, in particular, expression plasmids, for transferring genes into mammalian cells relies on routine techniques in the field of molecular biology. A basic text disclosing the general methods of preparing expression plasmids used in this invention is Molecular Cloning, A Laboratory Manual, 2nd Edition, eds. Sambrook et al., Cold Spring Harbor Laboratory Press, (1989), particularly Chapter 16 on Expression of Cloned Genes in Cultured Mammalian Cells. Examples 15C-D below set forth particular methods for preparing the expression plasmids used in the present invention. The particular vector used to transfer the antisense oligonucleotides of the present invention is not critical, and such vectors may include vectors derived from lambda and related phages or from filamentous phages. It is only necessary that the transferred nucleotide sequence encoding the antisense oligonucleotides of the present invention be expressed in the transfected tumor cell under conditions sufficient to reduce bcl-2 gene expression in the tumor cell. The present invention includes expression of the antisense oligonucleotide either from an extrachromosomal position (e.g. from an expression plasmid) or from a position integrated into the host genome itself, as mediated by other vectors, such as recombinant retroviral vectors (Reed et al. bcl-2 mediated tumorigenicity in a T-cell lymphoid cell line: synergy with C-MYC and inhibition by bcl-2 antisense. PNAS USA 87:3660 (1990)).

- 30 A. Treatment of Lymphoma Cells With 18-mer Synthetic bcl-2 Antisense Oligodeoxynucleotides.
 Lymphoma cell line SU-DHL-4, obtained from a use of diffuse, histiocytic, non-Hodgins lymphoma (Epstein et al. Two new monoclonal antibodies (LN-1, LN-2) reactive in B5 formalin-fixed, paraffin-embedded

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tissues with follicular center and mantle zone human B lymphocytes and derived tumors. J. Immunol. 133:1028 . (1984)) and containing a t(14;18) translocation was treated with 18-mer synthetic bcl-2-AS
5 oligodeoxynucleotides targeted for binding with the first six codons of the bcl-2 mRNA. As a control, SU-DHL-4 cells were treated with various control oligomers, including 18-mers having the same nucleoside composition as the AS oligomer, but in which the bases were in
10 scrambled order (SC) .

Aliquots of 1.5 ml of HL-1 serum-free medium (Ventrex Labs, Inc.) supplemented with 1 mM L-glutamine, 50 Units/ml penicillin, and 100 ug/ml streptomycin and either 5 ug of purified oligonucleotides or 30 ug of
15 Lipofectin® [1:1 w/w mixture of N-(1-2,3-dioleyloxy)propyl)-n,n,n-trimethylammonium chloride (DOTMA) and dioleoylphosphatidylethanolamine (DOPE)] were combined and added to 0.75×10^6 SU-DHL-4 cells in 3 mls of HL-1 medium. Cells were then either
20 cultured at 37°C in a humidified atmosphere of 5% CO₂/95% air in 24 well plates (2 mls/well) for immunoblot and RT-PCR assays, or in 96-well flat-bottom microtiter plates (0.1 ml/well) for MTT assays. For cells in microtiter cultures, typically 0.1 ml of additional HL-1
25 media with or without various chemotherapeutic drugs was added after 1 day, and the cells were cultured for an additional 2 days before performing MTT assays.

Cells were washed once in PBS, lysed in a buffer containing 1% Triton X100, and samples normalized for protein content (25 ug) prior to size-fractionation of proteins by SDS-PAGE (12% gels) and transfer to nitrocellulose filters for immunoblot assays as described in Reed et al. Cancer Res. 51:6529 (1991). Preliminary experiments determined that aliquots of lysates containing 25 ug of total protein produced results in the
30
35

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linear range of the assay. Blots were first incubated with 0.1% (v.v) of a rabbit antiserum directed against a synthetic peptide corresponding to amino-acids (aa) 41-54 of the human Bcl-2 protein, as shown in SEQ ID NO. 21 (id) followed by 2.8 ug/ml biotinylated goat anti-rabbit IgG (Vector Labs, Inc.). Bands corresponding to p26-Bcl-2 were then visualized by color development using a Horseradish Peroxidase (HRP)-avidin-biotin complex reagent (Vector Labs, Inc) and 3,3'-diaminobenzidine (DAB). Stained blots were then incubated with a second anti-Bcl-2 antibody directed against aa 61-76 of the Bcl-2 protein (SEQ ID NO. 21) followed by 0.25 uCi/ml ¹²⁵I-protein A. Bcl-2 bands were excised from the blots and subjected to gamma-counting.

Despite the mitochondrial location of Bcl-2 protein, no difference in the rate of MTT dye reduction by mitochondrial enzymes was noted in cells that were identical except for their levels of p26-Bcl-2. These comparisons were made using pairs of exponentially growing lymphoid cell lines that differed only in that one line had been stably infected with a recombinant bcl-2 retrovirus and the other with the parental retroviral vector lacking a bcl-2 cDNA insert (Miyashita et al. Cancer Res. 52:5407 (1992); Blood 81:151 (1993)).

Anticod specific reductions in the relative levels of bcl-2 mRNA were detected within 1 day by a semi-quantitative reverse transcriptase polymerase chain reaction (RT-PCR) assay. See Figure 8A.

SU-DHL-4 cells were cultured with 0.83 ug/ml of oligomers complexed with 5 ug of cationic lipids (Lipofectin; BRL/Gibco, Inc.) per ml of serum-free media y(13,19). In Figure 8A, total RNA was isolated from cells after 1 day and relative levels of bcl-2 and glyceraldehyde 3'-phosphate dehydrogenase (GAPDH) mRNAs

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were assessed by RT-PCR assay as described in Kitada et al. Antisense R & D 3:157 (1993).

In Figure 8(b), SU-DHL-4 cells were cultured with pairs of either PS (squares) or PO/PS (circles) As- and Sc-oligonomers for 3 days. Relative levels of Bcl-2 protein were then measured using a quantitative immunoblot assay, as described above, and the data expressed as a percentage relative to cells treated with control Sc-oligonomers. The inset shows immunoblot results for p26-Bcl-2 and a p75 cross-reactive (CR) band in a case where As-PO/PS oligomer produced a 41% relative decrease in Bcl-2 protein levels. In Figure 8(c), 10^{-4} M Ara-C, MTX, or DEX was added 1 day after addition of PS (squares) or PO/PS (circles) oligomers to cultures of SU-DHL-4 cells, and MTT assays were performed on day 3. Data are presented as a % control relative to cells cultured with drugs in the absence of any oligomers, and represent the results of 9 of 10 consecutive experiments [in one experiment, the MTT assay failed]. Similar results were obtained when dye exclusion assays were used to assess cell survival rather than MTT assay [not shown].

Mean values for the data are indicated by horizontal lines. Statistical analysis of the data was by paired t-test (As versus Sc). Concentrations of As- and Sc-oligonomers (≈ 150 nM) were adjusted to maximize As effects while maintaining sequence specificity.

Variations in the amounts of starting RNA were controlled for by RT-PCR analysis using primers specific for GAPDH mRNA.

The long half-life of the bcl-2 protein (approximately 14 hours) may account for the AS-mediated reductions in bcl-2 proteins not being as dramatic as for

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reductions in bcl-2 mRNA, taking longer to achieve (about 3 days), and appearing more variable.

Figure 8(c) shows the composite results for 10 experiments where relative levels of bcl-2 protein were compared in SU-DHL-4 cells treated with AS or SC oligomers. AS-mediated reductions in bcl-2 protein levels ranged from as much as 66% to as little as 10%, with an average relative reduction of about 30%, compared to SU-DHL-4 cells that were treated in the identical manner with control oligomers. Levels of a variety of control mitochondrial proteins such as F₁-beta-ATPase and cytochrome C, which like bcl-2 are encoded by nuclear genes, were not adversely affected by AS-oligomers (not shown), indicating that the AS-mediated reductions in bcl-2 protein levels were specific. The insert in Figure 8(b), for example, shows a comparison of p26-Bcl-2 with a 78-kDa protein that cross reacts with one of the rabbit antisera employed for immunoblot assays, demonstrating a decrease in the levels of p26-bcl-2 but not p78 in the AS-treated cells relative to cells that received control SC-oligomers.

B. Effect of Treatment of SU-DHL-4 Cells with bcl-2 AS Oligomers on Cell Sensitivity to Cancer Chemotherapeutic Agents

This study was performed to determine whether treatment of SU-DHL-4 cells with bcl-2 AS-oligomers could increase their relative sensitivity to killing by the cancer chemotherapeutic agents Ara-C, MTX, and DEX, which are anticancer drugs.

Previous control studies demonstrated that bcl-2 AS oligomers had little or no effect on SU-DHL-4 cell growth and survival at least during the first three days of culture (Kitada et al. Antisense R & D 3:157 (1993)). AS-mediated reductions in bcl-2 protein levels in these

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lymphoma cells as well as in other cells do not typically accelerate the rate of cell death in cultures unless the cells are deprived of serum growth factors (Reed et al. Proc. Natl. Acad. Sci. USA 87:3660 (1990)).

5 In the present work, preliminary studies demonstrated that more than 90% of SU-DHL-4 cells survived treatment for 4 days with high dose (10^{-4}) Ara-C, MTX or DEX, presumably because of their high levels of bcl-2 protein (Not shown). At these concentrations,
10 however, all drugs induced essentially complete inhibition of SU-DHL-4 cell proliferation, consistent with bcl-2 converts drugs from cytotoxic to cytostatic. Comparisons of AS and SC oligomers demonstrated that bcl-
15 2 AS treatment markedly enhanced the sensitivity of these lymphoma cells to MTX and Ara-C, and to a lesser extent to DEX (Figure 8(c)).

Despite some variability in results, on average, the addition of bcl-2 AS oligomers to cultures of SU-DHL-4 cells treated with MTX or Ara-C resulted in
20 79-84% greater inhibition (reduction in viable cell numbers) than use of either drug alone ($P < 0.002$ for AS versus SC) in the absence of introducing the bcl-2 AS oligomers of the invention. Statistically significant results were obtained for DEX-treated SU-DHL-4 cells
25 ($P = 0.01$). The 20-30% reduction in viable cell numbers observed for control oligomer-treated cells could reflect a degree of sequence non-specificity, but was probably related to the use of cationic lipids to facilitate oligomer delivery into cells.

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C. Effect of Transfected Cells With Expression Plasmids Encoding Human bcl-2 Protein on Sensitivity to Chemotherapeutic Agents.

To further confirm the sequence specificity of bcl-2 AS oligomers for enhancing sensitivity to chemotherapeutic anticancer drugs, a study was conducted using an Interleukin-3 (IL-3)-dependent murine hemopoietic cell line 32D.C13 that had been stably transfected with expression plasmid encoding either the human bcl-2 protein or a viral homolog of bcl-2, BHRF-1, which has only 22% homology with bcl-2. 32D.C13 cells were obtained from Dr. Giovanni Rovera of the Wistar Institute, Philadelphia, PA.

Treatment of 32D cells with oligomer/cationic lipid complexes was as described above except that 50 Units/ml of murine recombinant IL-3 (rIL-3) was included in the HL-1 media, the initial cell density was 10^5 per ml, and replication-defective adenovirus dl312 (MOI=200) was added 30 minutes after exposure of cells to oligomers to facilitate exit of DNA from endosomes [Yoshimura K, et al. J Biol Chem. 268, 2300, (1993)].

32D cells that had been stable transfected with expression plasmids encoding either human p26-Bcl-2 or EBV p19-BHRF-1 (Takayama, S. et al. submitted) were cultured in medium (10^5 /ml) containing IL-3 and PO/PS oligomers for 3 days to achieve reductions in human Bcl-2 protein levels. The cells were then retreated with oligomers alone (C) or in combination with various concentrations of MTX and the relative number of viable cells assessed by MTT assay 2 days later. Data represent mean +/- standard deviation for triplicate determinations and are expressed as a % relative to cells that received no MTX. Statistical analysis of data for 10^{-6} to 10^{-4} M MTX was by a 2-way Analysis of Variables method (Finney, D.J. In Statistical Methods in Biological Assays, p. 72,

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1978 (3rd edition, Charles Griffin & Co., London). Comparable results were obtained with dye exclusion assays [not shown].

5 RNAs derived from the human bcl-2 construct in 32D-BCL-2 cells were a target for bcl-2 AS oligomers, whereas RNAs from the BHRF-1 expression plasmid are not. Thus the chemosensitivity to cytotoxic drugs of 32D.C13 cells expressing BHRF-1 should have been unaffected by the AS treatment.

10 Preliminary experiments demonstrated that upon withdrawal of IL-3 from 32D.C13 cells, levels of endogenous mouse bcl-2 protein declined and the cells underwent apoptosis. bcl-2 and BHRF-1 comparably supported the survival of 32D.C13 cells in the absence of 15 IL-3, and the proliferative rates of 32D.C13 cells containing high levels of these proteins were similar in the presence of IL-3, thus excluding these variables as explanations for any differences in chemosensitivity.

20 Figures 9(a) and (b) compares the sensitivity of 32D-BCL-2 and 32D-BHRF-1 cells to various concentrations of MTX. Treatment with bcl-2 AS-oligomers resulted in sequence-specific increases in the sensitivity of 32D-BCL-2 cells to inhibition by MTX at concentrations of 10^{-6} to 10^{-4} M ($P \leq 0.001$ for AS versus SC). In contrast, treatment with bcl-2 AS oligomers produced no significant difference in the sensitivity of 25 32D-BHRF-1 cells to MTX, relative to control SC-oligomers (Figures 9(a) and (b)). These data indicate that the effects of bcl-2 AS oligomers on chemosensitivity to 30 cytotoxic agents drugs are sequence specific. Furthermore, several other control oligomer, including bcl-2 sense, other scrambled sequences with the same nucleoside composition as AS, and oligomers with totally unrelated

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sequences all had comparatively little effect on the chemosensitivity of the cells (Not shown).

The findings above demonstrated that bcl-2 AS oligomers produced sequence specific reductions in bcl-2 mRNA and bcl-2 protein levels and that these events were associated with increased sensitivity to chemotherapeutic agents such as anticancer drugs. The portion of tumor cells killed by the chemotherapeutic agents was greater than the portion killed by the same amount of chemotherapeutic agents in the absence of introducing the bcl-2 AS oligomers of the invention.

D. Effect of Transfecting Cells With Expression Plasmids Encoding Human bcl-2 Protein on Sensitivity of Lymphoma Cells to Chemotherapeutic Agents.

A different strategy was employed to determine if AS-mediated reductions in bcl-2 gene expression could be achieved with an inducible bcl-2 AS expression plasmid that used a heavy metal responsive human metallothionein-IIA promoter in another translocation t(14;18)-containing lymphoma line, RS11846. RS11846 was obtained from Dr. Carlo Croce (Wistar Institute, Philadelphia, PA (Tsujimoto and Croce, Proc. Natl. Acad. Sci. USA 83:5214 (1986)).

To prepare the expression plamid, a 0.91 kbp bcl-2 cDNA (*ibid*) was subcloned in either antisense (AS) or sense (S) orientatino into a HindIII site downstream of a human metallothionein-IIA promoter in the plasmid pMEP-4 (Invitrogen, Inc.), which contains a hygromycin phosphotransferase gene and the EBNA-1 gene and origin of DNA replication from Epstein Varr Virus for high copy episomal maintenance.

RS11846 cells (5×10^6) in Dulbecco's phosphate buffered saline containing 30 ug of plasmid DNA were

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electroporated (1500 uF, 270 V/cm) using a Cellject Electroporation System from EquiBio, Inc. Cells were returned to their usual culture media (RPMI-L 1640 supplemented with 10% fetal bovine serum, 1 mM L-glutamine, 50 Units/ml penicillin, and 100 ug/ml streptomycin) at 2 X 10⁵ cells per ml and cultured for 2 days before seeding cells at 2X10⁵ per ml in media containing 200 ug/ml hygromycin. After 3 weeks of culture, the resulting bulk cell lines were passaged in successively higher concentrations of hygromycin in 200 ug/ml increments until the concentration reached 1 mg/ml (about 4 weeks).

Hygromycin-resistant RS11846 cell were cultured in RPMI/10% serum media containing 0.5 uM CdCl₂ and 3 days later immunoblot assays were performed using 25 ug protein/lane essentially as described in Tanaka S, et al. J. Biol. Chem. 268, 10920 (1993) and in Reed et al. Cancer Res. 51:6529 (1991)).

As summarized in Figures 10(a), (b)(1), (b)(2), and (b)(3), control ("C") and bcl-2-As ("As") plasmids were introduced into RS11846 cells and expression was induced with either 0.5 uM CdCl₂ or 50 uM ZnCl₂ for various times. As an additional control, RS11846 cells containing inducible plasmids with the bcl-2 cDNA in sense ("S") orientation were also analyzed. RS11846 cells were induced for 3 days and relative levels of Bcl-2 and F₁- β -ATPase proteins were assessed by immunoblot assay of Tanaka et al. J. Biol. Chem. 268:10920 (1993). In Figure 10(a), RS11846 cells were cultured at 10⁵ cells/ml in medium containing 0.5 uM CdCl₂ and 1 day later 10⁻⁷ M Ara-C or an equivalent volume of diluent control was added. Relative numbers of viable cells were estimated from MTT assays at various times and the mean +/- S.D. calculated for triplicate samples. In Figures 10(b)(1-3), RS11846 cells were cultured as in Figure

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10(a), except that various concentrations of Ara-C, MTX,
or DEX were added. Data represent mean +/- S.D. for
triplicate samples. Statistical calculations are by
2-way Analysis of Variables. DEX served as a negative
5 control here since RS11846 cells have lost glucocorticoid
receptors.

Preliminary experiments demonstrated that
RS11846 cells tolerated the addition of up to 0.5 microM
CdCl₂ or to microM ZnCl₂ to cultures for one week,
10 experiencing a slight decrease in growth rate but
essentially no decline in percentage cell viability (Not
shown).

In the absence of heavy metal induction, the
relative levels of bcl-2 protein in RS11846 cells
15 containing the control or bcl-2 AS plasmid were
comparable, as determined by immunoblot assays (Not
shown). When 0.5 uM CdCl₂ or uM ZnCl₂ was added,
reductions in bcl-2 protein became evident in the AS-
expressing cells at 2 days and maximal inhibition of 30-
20 40% was obtained at three to four days, relative to
control RS11846 cells.

Figure 10(a) shows an example of immunoblot
data derived from RS11846 cells after three days of
exposure to 0.5 mM CdCl₂, demonstrating reduced levels of
25 bcl-2 protein in the AS-plasmid containing cells compared
to RS11846 cells that harbored the control plasmid. The
relative levels of a control mitochondrial protein F₁-
beta-ATPase were comparable in all cell lines, consistent
with sequence-specific alterations in bcl-2 protein
30 levels.

When RS11846 cells containing either the
control or bcl-2-AS plasmids were cultured for various
times in 0.5uM CdCl₂ or 50 uM ZnCl₂, no significant

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difference in the growth rates of these two cell lines was observed (Figure 8(b)). Thus, AS-mediated reductions in *Bcl-2* protein levels by themselves did not impair RS11846 cell proliferation or survival.

5 Inclusion of low-dose Ara-C (10^{-7} M) in the cultures of control RS11846 cells resulted in only a slight decline in the net numbers of viable cells, presumably because of the high levels of *Bcl-2* protein found in these *t(14;18)*-containing lymphoma cells. In
10 contrast, addition of 10^{-7} M Ara-C to cultures of *bcl-2*-AS expressing RS11846 cells was markedly inhibitory (Figure 8(b)). Ara-C, however, had no effect on *bcl-2* AS-expressing RS11846 cells in the absence of heavy metal induction of the MT promoter, when directly compared with
15 RS11846 cells containing the control plasmid under the same conditions [not shown]. Figure 8(c) shows that the enhanced sensitivity to Ara-C observed for *bcl-2*-AS-expressing RS11846 cells occurred over a wide range of drug concentrations ($P<0.001$). Heavy-metal
20 induction of the *bcl-2*-AS expression plasmid also significantly increased the relative sensitivity of RS11846 lymphoma cells to MTX ($P<0.001$), but not to DEX. Glucocorticoid receptor binding assays demonstrated that RS11846 cells have lost receptors for these steroid
25 hormones [not shown], thus providing a specificity control showing that AS-mediated reductions in *bcl-2* protein levels are by themselves insufficient to impair the growth or survival of these lymphoma cells.

30 Using a plurality of anticode approaches, the present invention demonstrated that average reductions of 30-40% in the relative levels of *bcl-2* protein markedly enhanced the sensitivity of lymphoma cells, in particular, *t(14;18)*-containing lymphoma cell lines to chemotherapeutic agents such as conventional anticancer drugs. These examples demonstrated that introducing the
35

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claimed anticodc oligomers into tumor cells achieves a reduction of *bcl-2* expression and increases the chemosensitivity of ne plastic cells to chemotherapeutic agents or anticancer drugs.

5 Accordingly, the present invention achieved a method of killing tumor cells by introducing to tumor cells anticodc oligomers which reduce *bcl-2* gene expression or impair Bcl-2 protein function before contacting the cells with chemotherapeutic agents including anticancer drugs. The conventional anticancer drugs reduced the numbers of viable malignant cells, and the portion of tumor cells killed was greater than the portion which would have been killed by the same amount of drug in the absence of introducing the anticodc oligomers into the cells.

10

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Having thus disclosed exemplary embodiments of the present invention, it should be noted by those skilled in the art that this disclosure is exemplary only and that various other alternatives, adaptations, and 20 modifications may be made within the scope of the present invention. Accordingly, the present invention is not limited to the specific emobodiments as illustrated herein, but is only limited by the following claims.

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SEQUENCE LISTING

(1) GENERAL INFORMATION:

(i) APPLICANT: Reed, John

(ii) TITLE OF INVENTION: Regulation of bcl-2 Gene Expression

(iii) NUMBER OF SEQUENCES: 23

(iv) CORRESPONDENCE ADDRESS:

- (A) ADDRESSEE: Gray, Cary, Ames & Frye
- (B) STREET: 401 B Street, Suite 1700
- (C) CITY: San Diego
- (D) STATE: CA
- (E) COUNTRY: USA
- (F) ZIP: 92101-4297

(v) COMPUTER READABLE FORM:

- (A) MEDIUM TYPE: Diskette 3.5 inch, 1.44 Mb storage
- (B) COMPUTER: IBM PC compatible
- (C) OPERATING SYSTEM: PC-DOS/MS-DOS
- (D) SOFTWARE: PatentIn Release #1.0, Version #1.25

(vi) CURRENT APPLICATION DATA:

- (A) APPLICATION NUMBER: US
- (B) FILING DATE:
- (C) CLASSIFICATION:

(viii) ATTORNEY/AGENT INFORMATION:

- (A) NAME: Brotman, Harris F.
- (B) REGISTRATION NUMBER: 35461
- (C) REFERENCE/DOCKET NUMBER: P0041US0

(ix) TELECOMMUNICATION INFORMATION:

- (A) TELEPHONE: (619) 699-3630
- (B) TELEFAX: (619) 236-1048

(2) INFORMATION FOR SEQ ID NO:1:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 20 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

CCCTTCCTAC CGCGTGCAG

20

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(2) INFORMATION FOR SEQ ID NO:2:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 35 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: NO

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

CTTTTCCTCT GGGAAAGGATG GCGCACGCTG GGAGA

35

(2) INFORMATION FOR SEQ ID NO:3:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 20 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

CCTCCGACCC ATCCACGTAG

20

(2) INFORMATION FOR SEQ ID NO:4:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 33 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: NO

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

ACGGGGTACG GAGGCTGGGT AGGTGCATCT GGT

33

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(2) INFORMATION FOR SEQ ID NO:5:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 20 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

GTTGACGTCC TACGGAAACA

20

(2) INFORMATION FOR SEQ ID NO:6:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 36 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: NO

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

CCCCCAAATG CAGGATGCCT TTGTGGAACT GTACGG

36

(2) INFORMATION FOR SEQ ID NO:7:

- (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 20 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: single
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: NO

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

GGGAAGGATG GCGCACGCTG

20

(2) INFORMATION FOR SEQ ID NO:8:

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- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 17 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: singl
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

CGCGTGCAC CCTCTTG

17

(2) INFORMATION FOR SEQ ID NO:9:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 17 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

TACCGCGTGC GACCCTC

17

(2) INFORMATION FOR SEQ ID NO:10:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 17 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

TCCTACCGCG TGCGACC

17

(2) INFORMATION FOR SEQ ID NO:11:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 17 bas pairs
 - (B) TYPE: nucleic acid

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- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

CCTTCCTTACCC GCGTGCG

17

(2) INFORMATION FOR SEQ ID NO:12:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 17 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

GACCCTTCCTT ACCGCGT

17

(2) INFORMATION FOR SEQ ID NO:13:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 17 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:13:

GGAGACCCTT CCTACCG

17

(2) INFORMATION FOR SEQ ID NO:14:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 15 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

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(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

CGGGCGGCAG CGCGG

15

(2) INFORMATION FOR SEQ ID NO:15:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 15 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:15:

CGGCGGGGGCG ACGGA

15

(2) INFORMATION FOR SEQ ID NO:16:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 16 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(iv) ANTI-SENSE: YES

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

CGGGAGCGCG GCGGGC

16

(2) INFORMATION FOR SEQ ID NO:17:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 18 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

TCTCCCAGCG TGCGCCAT

18

(2) INFORMATION FOR SEQ ID NO:18:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 18 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

TGCACTCACG CTCGGCCT

18

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(3) INFORMATION FOR SEQ ID NO:19:

- (i) SEQUENCE CHARACTERISTICS:
- (A) LENGTH: 5086 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

GCGCCCGCCC	CTCCGCGCCG	CCTGCCGCC	CGCCCGCCGC	GCTCCCGCCC	GCCGCTCTCC	60
GTGGCCCCGC	CGCGCTGCCG	CCGCCGCCGC	TGCCAGCGAA	GGTGCCGGGG	CTCCGGGCC	120
TCCCTGCCGG	CGGCCGTCAG	CGCTCGGAGC	GAACTGCGCG	ACGGGAGGTC	CGGGAGGCGA	180
CCGTAGTCGC	GCCGCCGCGC	AGGACCAGGA	GGAGGAGAAA	GGGTGCCAG	CCCGGAGGCG	240
GGGTGCGCCG	GTGGGGTGCA	GCGGAAGAGG	GGGTCCAGGG	GGGAGAACTT	CGTAGCAGTC	300
ATCCTTTTA	GGAAAAGAGG	GAAAAAATAA	AACCCTCCCC	CACCACCTCC	TTCTCCCCAC	360
CCCTCGCCGC	ACCACACACA	GCGCGGGCTT	CTAGCGCTCG	GCACCGGC	GCCAGGCGCG	420
TCCTGCCTTC	ATTATCCAG	CAGCTTTCG	GAAAATGCAT	TTGCTGTTCG	GAGTTAAC	480
AGAAGACGAT	TCCTGCCTCC	GTCCCCGGCT	CCTTCATCGT	CCCATCTCCC	CTGTCTCT	540
CCTGGGGAGG	CGTGAAGCGG	TCCC GTGGAT	AGAGATTCA	GCCTGTGTCC	GCGCGTGTGT	600
GCGCGCGTAT	AAATTGCCGA	GAAGGGAAA	ACATCACAGG	ACTTCTGCGA	ATACCGGACT	660
GAAAATTGTA	ATTCACTCTGC	CGCCGCCGCT	GCCAAAAAAA	AACTCGAGCT	CTTGAGATCT	720
CCGGTTGGGA	TTCCCTGCCA	TTGACATTT	TGTGAAGCAG	AAGTCTGGGA	ATCGATCTGG	780
AAATCCTCCT	AATTTTACT	CCCTCTCCCC	CCGACTCCTG	ATTCAATTGG	AAGTTCAA	840
TCAGCTATAA	CTGGAGAGTG	CTGAAGATTG	ATGGGATCGT	TGCCTTATGC	ATTTGTTTG	900
GTTTTACAAA	AAGGAAACTT	GACAGAGGAT	CATGCTGTAC	TTAAAAAATA	CAAGTAAGTC	960
TCGCACAGGA	AATTGGTTTA	ATGTAAC	CAATGGAAAC	CTTGAGATT	TTTTACTTAA	1020
AGTGCATTG	AGTAAATT	ATTCCAGGC	AGCTTAATAC	ATTGTTTTA	GCCGTGTTAC	1080
TTGTAGTGTG	TATGCCCTGC	TTTCACTCAG	TGTGTACAGG	GAAACGCACC	TGATTTTTA	1140
CTTATTAGTT	TGTTTTCT	TTAACCTTTC	AGCATCACAG	AGGAAGTAGA	CTGATATTAA	1200
CAATACTTAC	TAATAATAAC	GTGCCTCATG	AAATAAAGAT	CCGAAAGGAA	TTGGAATAAA	1260
AATTTCCTGC	GTCTCATGCC	AAGAGGGAAA	CACCAGAAC	AAGTGTCCG	CGTGATTGAA	1320

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GACACCCCT CGTCCAAGAA	TGCAAAGCAC	ATCCAATAAA	ATAGCTGGAT	TATAACTCCT	1380
CTTCTTCCTC TGGGGGCCGT	GGGGTGGGAG	CTGGGGCGAG	AGGTGCCGTT	GGCCCCCGTT	1440
GCTTTCCCTC TGGGAAGGAT	GGCGCACGCT	GGGAGAACGG	GGTACGACAA	CCGGGAGATA	1500
GTGATGAAGT ACATCCATTA	TAAGCTGTCG	CAGAGGGCT	ACGAGTGGGA	TGCGGGAGAT	1560
GTGGCGCCG CGCCCCCGGG	GGCCGCCCCC	GCACCGGGCA	TCTTCTCCTC	CCAGCCCCGG	1620
CACACGCCCC ATCCAGCCGC	ATCCCACGAC	CCGGTCGCCA	GGACCTCGCC	GCTGCAGACC	1680
CCGGCTGCC CCGCGGCCGC	CGCGGGGCCT	GCGCTCAGCC	CGGTGCCACC	TGTGGTCCAC	1740
CTGGCCCTCC GCCAAGCCGG	CGACGACTTC	TCCCACGCT	ACCGCGGCAG	CTTCGCGGAG	1800
ATGTCCAGCC AGCTGCACCT	GACGCCCTTC	ACCGCGCGGG	GACGCTTGC	CACGGTGGTG	1860
GAGGAGCTCT TCAGGGACGG	GGTGAACCTGG	GGGAGGATTG	TGGCCTTCTT	TGAGTTCGGT	1920
GGGGTCATGT GTGTGGAGAG	CGTCAACCGG	GAGATGTCGC	CCCTGGTGGA	CAACATCGCC	1980
CTGTGGATGA CTGAGTACCT	GAACCGGCAC	CTGCACACCT	GGATCCAGGA	TAACGGAGGC	2040
TGGGATGCCT TTGTGGAAC	GTACGGCCCC	AGCATGCGGC	CTCTGTTGA	TTTCTCCTGG	2100
CTGTCTCTGA AGACTCTGCT	CAGTTGGCC	CTGGTGGGAG	CTTGCATCAC	CCTGGGTGCC	2160
TATCTGAGCC ACAAGTGAAG	TCAACATGCC	TGCCCCAAC	AAATATGCAA	AAGGTTCACT	2220
AAAGCAGTAG AAATAATATG	CATTGTCAGT	GATGTACCAT	GAAACAAAGC	TGCAGGCTGT	2280
TTAAGAAAAA ATAACACACA	TATAAACATC	ACACACACAG	ACAGACACAC	ACACACACAA	2340
CAATTAACAG TCTTCAGGCA	AAACGTCGAA	TCAGCTATTT	ACTGCCAAAG	GGAAATATCA	2400
TTTATTTTTT ACATTATTAA	AAAAAAAGAT	TTATTTATTT	AAGACAGTCC	CATCAAAACT	2460
CCGTCTTGG AAATCCGACC	ACTAATTGCC	AAACACCGCT	TCGTGTGGCT	CCACCTGGAT	2520
GTTCTGTGCC TGAAACATA	GATTGCTTT	CCATGTTGTT	GGCCGGATCA	CCATCTGAAG	2580
AGCAGACGGA TGGAAAAGG	ACCTGATCAT	TGGGAAAGCT	GGCTTCTGG	CTGCTGGAGG	2640
CTGGGGAGAA GGTGTTCA	CACTTGCATT	TCTTGCCT	GGGGCGTGA	TATTAACAGA	2700
GGGAGGGTTC CGTGGGGGG	AACTCCATGC	CTCCCTGGCC	TGAAGAAGAG	ACTCTTGCA	2760
TATGACTCAC ATGATGCATA	CCTGGTGGGA	GGAAAAGAGT	TGGGAACTTC	AGATGGACCT	2820
AGTACCCACT GAGATTCCA	CGCCGAAGGA	CAGCGATGGG	AAAAATGCC	TTAAATCATA	2880
GGAAAGTATT TTTTAAGCT	ACCAATTGTG	CCGAGAAAAG	CATTTAGCA	ATTTATACAA	2940
TATCATCCAG TACCTTAAAC	CCTGATTGTG	TATATTACATA	TATTTGGAT	ACGCACCCCC	3000

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CAACTCCCAA TACTGGCTCT GTCTGAGTAA GAAACAGAAT CCTCTGGAAC TTGAGGAAGT	3060
GAACATTTCG GTGACTTCCG ATCAGGAAGG CTAGAGTTAC CCAGAGCATC AGGCCGCCAC	3120
AAGTGCCTGC TTTTAGGAGA CCGAAGTCCG CAGAACCTAC CTGTGTCCC GCTTGGAGGC	3180
CTGGTCCTGG AACTGAGCCG GGCCCTCACT GGCCCTCTCC AGGGATGATC AACAGGGTAG	3240
TGTGGTCTCC GAATGTCTGG AAGCTGATGG ATGGAGCTCA GAATTCCACT GTCAAGAAAG	3300
AGCAGTAGAG GGGTGTGGCT GGGCCTGTCA CCCTGGGCC CTCCAGGTAG GCCCGTTTC	3360
ACGTGGAGCA TAGGAGCCAC GACCCTTCTT AAGACATGTA TCACTGTAGA GGGAGGAAC	3420
AGAGGCCCTG GGCCTTCCTA TCAGAAGGAC ATGGTGAAGG CTGGAACGT GAGGAGAGGC	3480
AATGGCCACG GCCCATTTC GCTGTAGCAC ATGGCACGTT GGCTGTGTGG CCTTGGCCAC	3540
CTGTGAGTTT AAAGCAAGGC TTTAAATGAC TTTGGAGAGG GTCACAAATC CTAAAAGAAG	3600
CATTGAAGTG AGGTGTCATG GATTAATTGA CCCCTGTCTA TGGAATTACA TGTAAAACAT	3660
TATCTTGTCA CTGTAGTTG GTTTTATTTG AAAACCTGAC AAAAAAAAAG TTCCAGGTGT	3720
GGAATATGGG GGTTATCTGT ACATCCTGGG GCATTAAAAA AAAATCAATG GTGGGAACT	3780
ATAAAAGAAGT AACAAAAGAA GTGACATCTT CAGCAAATAA ACTAGGAAAT TTTTTTTCT	3840
TCCAGTTAG AATCAGCCTT GAAACATTGA TGGAATAACT CTGTGGCATT ATTGCATTAT	3900
ATACCATTAA TCTGTATTAA CTTTCCAATG TACTCTGTTA AATGTTAAT GCTGTGGTTG	3960
ATATTCGAA AGCTGCTTTA AAAAAATACA TGCATCTCAG CGTTTTTTG TTTTAATTG	4020
TATTTAGTTA TGGCCTATAC ACTATTTGTG AGCAAAGGTG ATCGTTTCT GTTTGAGATT	4080
TTTATCTCTT GATTCTCAA AAGCATTCTG AGAAGGTGAG ATAAGCCCTG AGTCTCAGCT	4140
ACCTAAGAAA AACCTGGATG TCACTGGCCA CTGAGGAGCT TTGTTCAAC CAAGTCATGT	4200
GCATTTCCAC GTCAACAGAA TTGTTTATTG TGACAGTTAT ATCTGTTGTC CCTTTGACCT	4260
TGTTTCTTGA AGGTTCCCTC GTCCCTGGC AATTCCGCAT TTAATTCATG GTATTCAAGGA	4320
TTACATGCAT GTTTGGTTAA ACCCATGAGA TTCATTCACT TAAAAATCCA GATGGCGAAT	4380
GACCAGCAGA TTCAAATCTA TGGTGGTTTG ACCTTAGAG AGTTGCTTTA CGTGGCCTGT	4440
TTCAACACAG ACCCACCCAG AGCCCTCCTG CCCTCCTTCC GCGGGGGCTT TCTCATGGCT	4500
GTCCTTCAGG GTCTTCCCTGA AATGCAGTGG TCGTTACGCT CCACCAAGAA AGCAGGAAAC	4560
CTGTGGTATG AAGCCAGACC TCCCCGGCGG GCCTCAGGGA ACAGAATGAT CAGACCTTG	4620
AATGATTCTA ATTTTAAGC AAAATATTAT TTTATGAAAG GTTTACATTG TCAAAGTGAT	4680

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GAATATGGAA TATCCAATCC TGTGCTGCTA TCCTGCCAAA ATCATTAA TGGAGTCAGT	4740
TTGCAGTATG CTCCACGTGG TAAGATCCTC CAAGCTGCTT TAGAAGTAAC AATGAAGAAC	4800
GTGGACGTTT TTAATATAAA GCCTGTTTG TCTTTGTTG TTGTTCAAAC GGGATTCAACA	4860
GAGTATTGAA AAAATGTATA TATATTAAGA GGTCACGGGG GCTAATTGCT AGCTGGCTGC	4920
CTTTGCTGT GGGGTTTGT TACCTGGTTT TAATAACAGT AAATGTGCC AGCCTCTTGG	4980
CCCCAGAACT GTACAGTATT GTGGCTGCAC TTGCTCTAAG AGTAGTTGAT GTTGCATTTT	5040
CCTTATTGTT AAAAACATGT TAGAAGCAAT GAATGTATAT AAAAGC	5086

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(2) INFORMATION FOR SEQ ID NQ:20:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 717 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 1..717

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:

ATG GCG CAC GCT GGG AGA ACG GGG TAC GAC AAC CCG GAG ATA GTG ATG Met Ala His Ala Gly Arg Thr Gly Tyr Asp Asn Arg Glu Ile Val Met	48
1 5 10 15	15
AAG TAC ATC CAT TAT AAG CTG TCG CAG AGG GGC TAC GAG TGG GAT GCG Lys Tyr Ile His Tyr Lys Leu Ser Gln Arg Gly Tyr Glu Trp Asp Ala	96
20 25 30	
GGA GAT GTG GGC GCC GCG CCC CCG GGG GCC GCC CCC GCA CCG GGC ATC Gly Asp Val Gly Ala Ala Pro Pro Gly Ala Ala Pro Ala Pro Gly Ile	144
35 40 45	
TTC TCC TCC CAG CCC GGG CAC ACG CCC CAT CCA GCC GCA TCC CGC GAC Phe Ser Ser Gln Pro Gly His Thr Pro His Pro Ala Ala Ser Arg Asp	192
50 55 60	
CCG GTC GCC AGG ACC TCG CCG CTG CAG ACC CCG GCT GCC CCC GGC GCC Pro Val Ala Arg Thr Ser Pro Leu Gln Thr Pro Ala Ala Pro Gly Ala	240
65 70 75 80	
GCC GCG GGG CCT GCG CTC AGC CCG GTG CCA CCT GTG GTC CAC CTG GCC Ala Ala Gly Pro Ala Leu Ser Pro Val Pro Pro Val Val His Leu Ala	288
85 90 95	
CTC CGC CAA GCC GGC GAC GAC TTC TCC CGC CGC TAC CGC GGC GAC TTC Leu Arg Gln Ala Gly Asp Asp Phe Ser Arg Arg Tyr Arg Gly Asp Phe	336
100 105 110	
GCC GAG ATG TCC AGC CAG CTG CAC CTG ACG CCC TTC ACC GCG CGG GGA Ala Glu Met Ser Ser Gln Leu His Leu Thr Pro Phe Thr Ala Arg Gly	384
115 120 125	
CGC TTT GCC ACG GTG GTG GAG GAG CTC TTC AGG GAC GGG GTG AAC TGG Arg Phe Ala Thr Val Val Glu Glu Leu Phe Arg Asp Gly Val Asn Trp	432
130 135 140	

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GGG AGG ATT GTG GCC TTC TTT GAG TTC GGT GGG GTC ATG TGT GTG GAG Gly Arg Ile Val Ala Phe Phe Glu Phe Gly Gly Val Met Cys Val Glu 145 150 155 160	480
AGC GTC AAC CGG GAG ATG TCG CCC CTG GTG GAC AAC ATC GCC CTG TGG Ser Val Asn Arg Glu Met Ser Pro Leu Val Asp Asn Ile Ala Leu Trp 165 170 175	528
ATG ACT GAG TAC CTG AAC CGG CAC CTG CAC ACC TGG ATC CAG GAT AAC Met Thr Glu Tyr Leu Asn Arg His Leu His Thr Trp Ile Gln Asp Asn 180 185 190	576
GGA GGC TGG GAT GCC TTT GTG GAA CTG TAC GGC CCC AGC ATG CGG CCT Gly Gly Trp Asp Ala Phe Val Glu Leu Tyr Gly Pro Ser Met Arg Pro 195 200 205	624
CTG TTT GAT TTC TCC TGG CTG TCT CTG AAG ACT CTG CTC AGT TTG GCC L u Phe Asp Phe Ser Trp Leu Ser Leu Lys Thr Leu Leu Ser Leu Ala 210 215 220	672
CTG GTG GGA GCT TGC ATC ACC CTG GGT GCC TAT CTG AGC CAC AAG Leu Val Gly Ala Cys Ile Thr Leu Gly Ala Tyr Leu Ser His Lys 225 230 235	717

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(2) INFORMATION FOR SEQ ID NO:21: .

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 239 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

Met Ala His Ala Gly Arg Thr Gly Tyr Asp Asn Arg Glu Ile Val Met
1 5 10 15

Lys Tyr Ile His Tyr Lys Leu Ser Gln Arg Gly Tyr Glu Trp Asp Ala
20 25 30

Gly Asp Val Gly Ala Ala Pro Pro Gly Ala Ala Pro Ala Pro Gly Ile
35 40 45

Phe Ser Ser Gln Pro Gly His Thr Pro His Pro Ala Ala Ser Arg Asp
50 55 60

Pro Val Ala Arg Thr Ser Pro Leu Gln Thr Pro Ala Ala Pro Gly Ala
65 70 75 80

Ala Ala Gly Pro Ala Leu Ser Pro Val Pro Pro Val Val His Leu Ala
85 90 95

Leu Arg Gln Ala Gly Asp Asp Phe Ser Arg Arg Tyr Arg Gly Asp Phe
100 105 110

Ala Glu Met Ser Ser Gln Leu His Leu Thr Pro Phe Thr Ala Arg Gly
115 120 125

Arg Phe Ala Thr Val Val Glu Glu Leu Phe Arg Asp Gly Val Asn Trp
130 135 140

Gly Arg Ile Val Ala Phe Phe Glu Phe Gly Gly Val Met Cys Val Glu
145 150 155 160

Ser Val Asn Arg Glu Met Ser Pro Leu Val Asp Asn Ile Ala Leu Trp
165 170 175

Met Thr Glu Tyr Leu Asn Arg His Leu His Thr Trp Ile Gln Asp Asn
180 185 190

Gly Gly Trp Asp Ala Phe Val Glu Leu Tyr Gly Pro Ser Met Arg Pro
195 200 205

Leu Phe Asp Phe Ser Trp Leu Ser Leu Lys Thr Leu Leu Ser Leu Ala
210 215 220

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Leu Val Gly Ala Cys Ile Thr Leu Gly Ala Tyr Leu Ser His Lys
 225 230 235 •

(2) INFORMATION FOR SEQ ID NO:22:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 615 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: DNA (genomic)

(ix) FEATURE:

- (A) NAME/KEY: CDS
- (B) LOCATION: 1..615

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

ATG GCG CAC GCT GGG AGA ACG GGG TAC GAC AAC CGG GAG ATA GTG ATG	48
Met Ala His Ala Gly Arg Thr Gly Tyr Asp Asn Arg Glu Ile Val Met	
1 5 10 15	
AAG TAC ATC CAT TAT AAG CTG TCG CAG AGG GGC TAC GAG TGG GAT GCG	96
Lys Tyr Ile His Tyr Lys Leu Ser Gln Arg Gly Tyr Glu Trp Asp Ala	
20 25 30	
GGA GAT GTG GGC GCC GCG CCC CCG GGG GCC GCC CCC GCA CCG GGC ATC	144
Gly Asp Val Gly Ala Ala Pro Pro Gly Ala Ala Pro Ala Pro Gly Ile	
35 40 45	
TTC TCC TCC CAG CCC GGG CAC ACG CCC CAT CCA GCC GCA TCC CGC GAC	192
Phe Ser Ser Gln Pro Gly His Thr Pro His Pro Ala Ala Ser Arg Asp	
50 55 60	
CCG GTC GCC AGG ACC TCG CCG CTG CAG ACC CCG GCT GCC CCC GGC GCC	240
Pro Val Ala Arg Thr Ser Pro Leu Gln Thr Pro Ala Ala Pro Gly Ala	
65 70 75 80	
GCC GCG GGG CCT GCG CTC AGC CCG GTG CCA CCT GTG GTC CAC CTG GCC	288
Ala Ala Gly Pro Ala Leu Ser Pro Val Pro Pro Val Val His Leu Ala	
85 90 95	

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CTC CGC CAA GCC GGC GAC GAC TTC TCC CGC CGC TAC CGC GGC GAC TTC Leu Arg Gln Ala Gly Asp Asp Phe Ser Arg Arg Tyr Arg Gly Asp Phe	336
100 105 110	
GCC GAG ATG TCC AGC CAG CTG CAC CTG ACG CCC TTC ACC GCG CGG GGA Ala Glu Met Ser Ser Gln Leu His Leu Thr Pro Phe Thr Ala Arg Gly	384
115 120 125	
CGC TTT GCC ACG GTG GTG GAG GAG CTC TTC AGG GAC GGG GTG AAC TGG Arg Phe Ala Thr Val Val Glu Glu Leu Phe Arg Asp Gly Val Asn Trp	432
130 135 140	
GGG AGG ATT GTG GCC TTC TTT GAG TTC GGT GGG GTC ATG TGT GTG GAG Gly Arg Ile Val Ala Phe Phe Glu Phe Gly Gly Val Met Cys Val Glu	480
145 150 155 160	
AGC GTC AAC CGG GAG ATG TCG CCC CTG GTG GAC AAC ATC GCC CTG TGG Ser Val Asn Arg Glu Met Ser Pro Leu Val Asp Asn Ile Ala Leu Trp	528
165 170 175	
ATG ACT GAG TAC CTG AAC CGG CAC CTG CAC ACC TGG ATC CAG GAT AAC Met Thr Glu Tyr Leu Asn Arg His Leu His Thr Trp Ile Gln Asp Asn	576
180 185 190	
GGA GGC TGG GTA GGT GCA TCT GGT GAT GTG AGT CTG GGC Gly Gly Trp Val Gly Ala Ser Gly Asp Val Ser Leu Gly	615
195 200 205	

(2) INFORMATION FOR SEQ ID NO:23:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 205 amino acids
 - (B) TYPE: amino acid
 - (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

Met Ala His Ala Gly Arg Thr Gly Tyr Asp Asn Arg Glu Ile Val Met	
1 5 10 15	
Lys Tyr Ile His Tyr Lys Leu Ser Gln Arg Gly Tyr Glu Trp Asp Ala	
20 25 30	
Gly Asp Val Gly Ala Ala Pro Pro Gly Ala Ala Pro Ala Pro Gly Ile	
35 40 45	
Phe Ser Ser Gln Pro Gly His Thr Pro His Pro Ala Ala Ser Arg Asp	
50 55 60	
Pro Val Ala Arg Thr Ser Pro Leu Gln Thr Pro Ala Ala Pro Gly Ala	
65 70 75 80	

-74-

Ala Ala Gly Pro Ala Leu Ser Pro Val Pro Pro Val Val His Leu Ala
85 90 95

Leu Arg Gln Ala Gly Asp Asp Phe Ser Arg Arg Tyr Arg Gly Asp Phe
100 105 110

Ala Glu Met Ser Ser Gln Leu His Leu Thr Pro Phe Thr Ala Arg Gly
115 120 125

Arg Phe Ala Thr Val Val Glu Glu Leu Phe Arg Asp Gly Val Asn Trp
130 135 140

Gly Arg Ile Val Ala Phe Phe Glu Phe Gly Gly Val Met Cys Val Glu
145 150 155 160

Ser Val Asn Arg Glu Met Ser Pro Leu Val Asp Asn Ile Ala Leu Trp
165 170 175

Met Thr Glu Tyr Leu Asn Arg His Leu His Thr Trp Ile Gln Asp Asn
180 185 190

Gly Gly Trp Val Gly Ala Ser Gly Asp Val Ser Leu Gly
195 200 205

CLAIMS:

1. An anticode oligomer which binds to a site in mRNA transcribed from the human bcl-2 gen , said site being a strategic site selected from the splice donor site, the splice acceptor site and the or a portion of the 5'-untranslated region of said mRNA, which anticode oligomer, when brought into contact with cancer cells expressing the human bcl-2 gene, has the propefty of reducing the expression of at least one bcl-2 gene product and thereby inducing programmed cell death of said cancer cells.

2. An anticode oligomer as claimed in claim 1 wherein said anticode oligomer is an antisense oligonucleotide or analogue thereof.

3. An anticode oligomer as claimed in claim 1 or claim 2 wherein said anticode oligomer is substantially complementary to said strategic site.

4. An anticode oligomer as claimed in any one of the preceding claims wherein said strategic site is in the 5'-untranslated region of said mRNA expressed from the human bcl-2 gene and wherein said anticode oligomer binds exclusively to the or a portion of the 5'-untranslated region.

5. An anticode oligomer as claimed in any one of claims 2 to 4 wherein said anticode oligomer is a phosphodiester/methylphosphonate chimera or a phosphodiester/phosphorothioate chimera or a methylphosphonate/phosphorothioate chimera.

6. An anticode oligomer as claimed in any one of claims 2 to 4 wherein said anticode oligomer or chirally enriched form thereof is comprised of one or more groups selected from methylphosphonates,

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phosphorothioates, diesters, phosphorodithioates, or 2' substituted derivatives.

7. An anticode oligomer as claimed in any one of the preceding claims which is two to two hundred nucleotide bases long.

8. An anticode oligomer as claimed in any one of the preceding claims which is ten to forty nucleotide bases long.

9. An anticode oligomer as claimed in any one of the preceding claims wherein the cells expressing the human bcl-2 gene are human tumor cells.

10. An anticode oligomer as claimed in claim 8 wherein the strategic site is a sequence portion in pre-mRNA expressed from said bcl-2 gene.

11. An anticode oligomer as claimed in claim 10 wherein said anticode oligomer is substantially complementary to and binds to a strategic site in said pre-mRNA.

12. An anticode oligomer as claimed in claim 10 or claim 11 wherein said strategic site in said pre-mRNA is selected from the translation initiation site, the splice donor site, the splice acceptor site, codons 1 to 6 of the bcl-2 open reading frame and the or a portion of the 5'-untranslated region.

13. An anticode oligomer as claimed in any one of claims 9 to 12 which is two to two hundred nucleotide bases long.

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14. An anticode oligomer as claimed in any one of claims 9-13 which is ten to forty nucleotide bases long.

5 15. An anticode oligomer as claimed in any one of claims 9 to 13 wherein the cells expressing the human bcl-2 gene are human tumor cells.

10 16. An anticode oligomer useful for inhibiting the growth of cells expressing the human bcl-2 gene which anticode oligomer binds with a strategic site of RNA expressed from the human bcl-2 gene, which anticode oligomer is a phosphodiester/methylphosphonate chimera or a phosphodiester/phosphorothioate chimera or a methylphosphonate/phosphorothioate chimera of an antisense oligonucleotide which when brought into contact 15 with cancer cells expressing the human bcl-2 gene, has the property of reducing the expression of at least one bcl-2 gene product and thereby inducing programmed cell death of said cancer cells.

20 17. An anticode oligomer useful for inhibiting the growth of cells expressing the human bcl-2 gene comprising an anticode oligomer which binds with a strategic site of RNA expressed from the human bcl-2 gene, which anticode oligomer or chirally enriched form thereof is comprised of one or more groups selected from 25 methylphosphonates, phosphorothioates, diesters, phosphorodithioates, or 2' substituted derivatives, which anticode oligomer when brought into contact with cancer cells expressing the human bcl-2 gene, has the property of reducing the expression of at least one bcl-2 gene product and thereby inducing programmed cell death of 30 said cancer cells.

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18. A vector for transfecting human cancer cells comprising a nucleotide sequence that encodes an anticode oligomer as claim d in any one of claims 1 to 4.

5 19. A vector for transfecting human cancer cells comprising a nucleotide sequence that encodes an anticode oligomer which reduces expression from the human bcl-2 gene in said cells, said vector being derived from a phage.

10 20. A composition useful for inhibiting the growth of cells expressing the human bcl-2 gene which comprises an anticode oligomer as claimed in any one of claims 1 to 17 together with a pharmaceutically acceptable carrier.

15 21. A composition useful for inhibiting the growth of cells expressing the human bcl-2 gene which comprises a vector as claimed in claim 18 or claim 19 and a pharmaceutically acceptable carrier.

20 22. A composition as claimed in claim 20 wherein said pharmaceutically acceptable carrier is a cationic lipid.

23. A composition as claimed in claim 20 wherein said anticode oligomer is encapsulated in liposomes.

25 24. Use of an anticode oligomer as claimed in any one of claims 1 to 17 in the manufacture of a medicament for treating human cancer.

25. Use of a vector as claimed in claim 18 or claim 19 in the manufacture of a medicament for treating human cancer.

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26. Use of an anticode oligomer which binds with a sequence portion of mRNA expressed from the human bcl-2 gene in the manufacture of a medicament for treating solid tumors in humans.

5 27. Use of an anticode oligomer which binds with a sequence portion of mRNA expressed from the human bcl-2 gene in the manufacture of a medicament for treating breast cancer.

10 28. Materials for simultaneous or sequential use for killing human cancer cells expressing the bcl-2 gene comprising:

an anticode oligomer which binds to a sequence portion of mRNA expressed from a human bcl-2 gene and;

15 a chemotherapeutic agent toxic to said human cancer cells.

20 29. Materials as claimed in claim 28 wherein said chemotherapeutic agent is selected from antimetabolites, alkylating agents, plant alkaloids and antibiotics.

30. Materials as claimed in claim 29 wherein said chemotherapeutic agent further comprises a radionuclide.

25 31. Materials as claimed in claim 28 or claim 29 wherein said anticode oligomer is as claimed in any one of claims 1 to 17.

32. A method of inducing programmed cell death of cancer cells which express the human bcl-2 gene comprising the steps of:

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- a. providing an antic de oligomer as claimed in any one of claims 1 to 17; and
- b. contacting said cells with said anticode oligomer under conditions sufficient to induce programmed cell death in said cells.

5

10

33. A method of killing human cancer cells which express the human bcl-2 gene comprising the steps of:

- a. providing an anticode oligomer which binds with a sequence portion of mRNA expressed from the human bcl-2 gene;
- b. contacting said cells with said anticode oligomer under conditions to inhibit growth of said cells; and
- c. contacting said cells with an amount of at least one secondary therapeutic agent toxic to said cells sufficient to kill at least a portion thereof.

15

20

34. A method as claimed in claim 33 wherein said anticode oligomer is as claimed in any one of claims 1 to 17.

25

35. A method as claimed in claim 33 wherein said secondary agent is a chemotherapeutic agent.

36. A method as claimed in any one of claims 33-35 wherein said chemotherapeutic agent is selected from antimetabolites, alkylating agents, plant alkaloids and antibiotics.

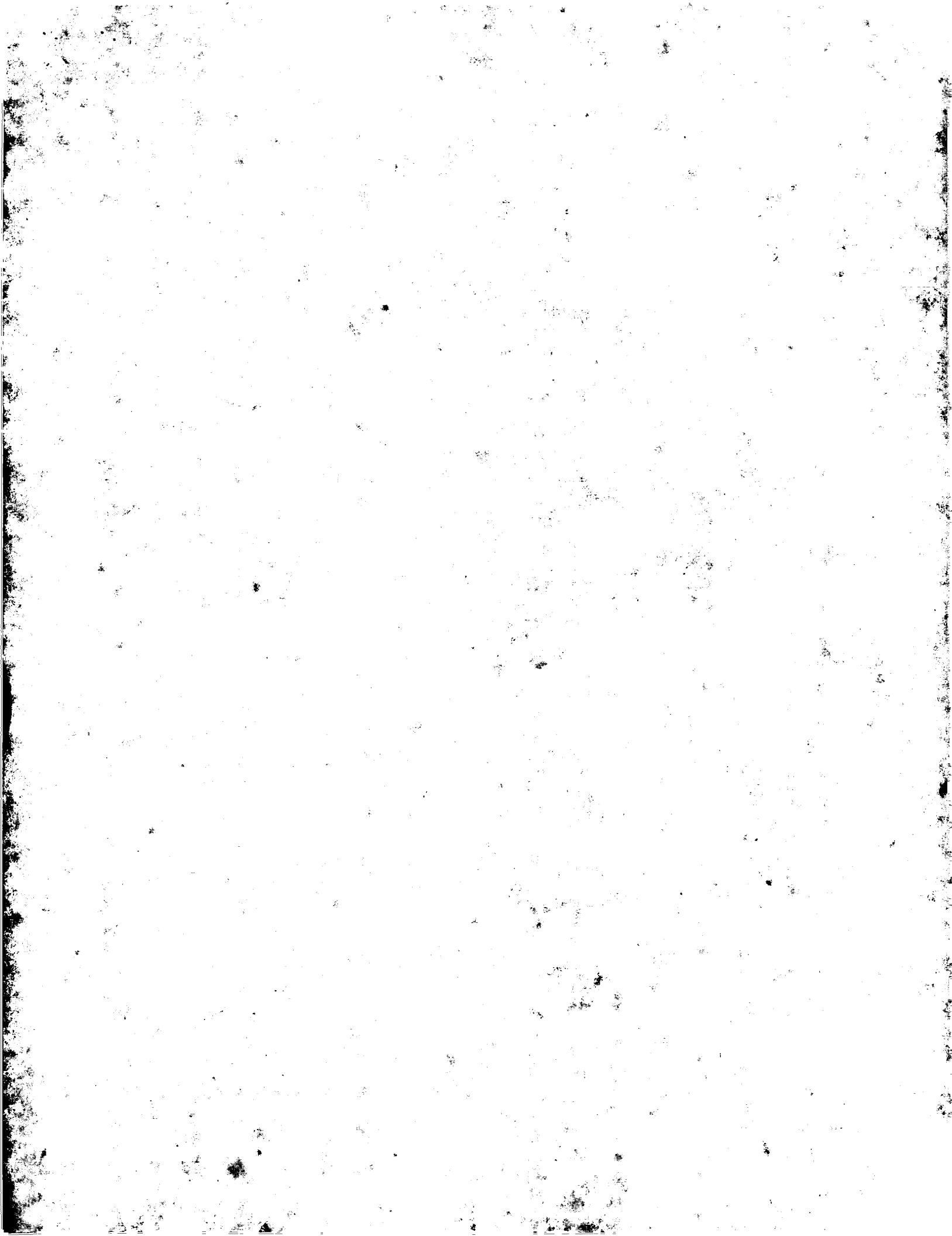
-81-

37. A method as claimed in claims 33 to 34 wherein said secondary therapeutic agent is a radiotherapeutic agent.

38. A method as claimed in claims 33, 34, or
5 37 in which said step of contacting said cells with a radiotherapeutic agent comprises contacting said cells with a beam of radiation, wherein said radiation comprises either particle radiation or non-particle radiation generated from a source which is non-contiguous
10 with said cells.

39. A method as claimed in claims 33, 34, or
37 in which said step of contacting said cells with a radiotherapeutic agent comprises contacting said cells with a radionuclide.

15 40. A composition useful for killing human cancer cells expressing the bcl-2 gene which comprises an anticode oligomer as claimed in any one of claims 1-17 and a chemotherapeutic agent toxic to said human cancer cells.



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FIG. 1 (a)

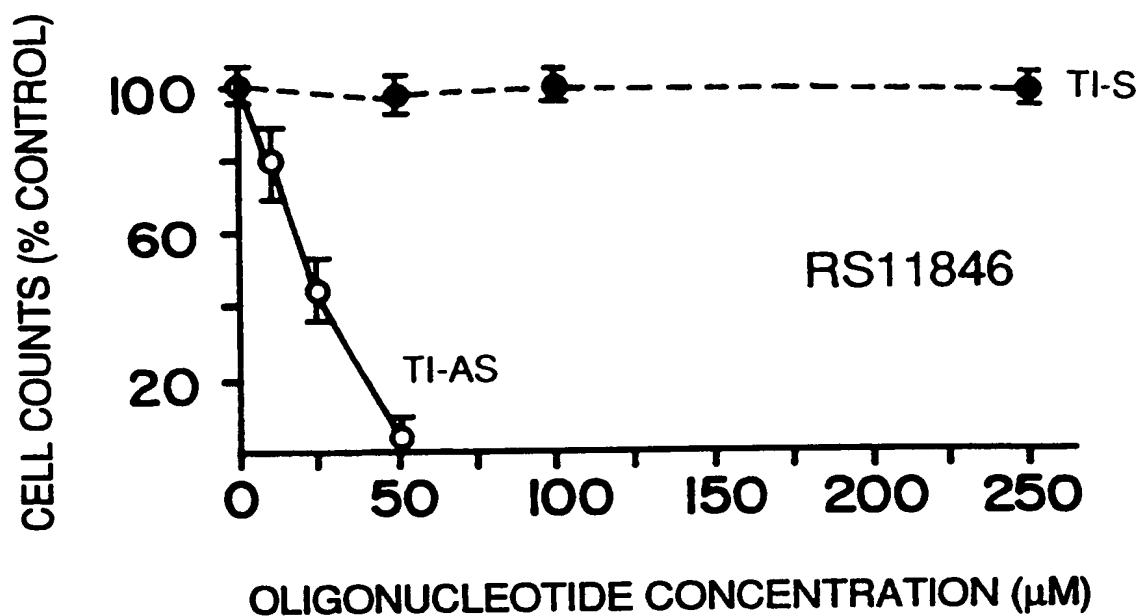
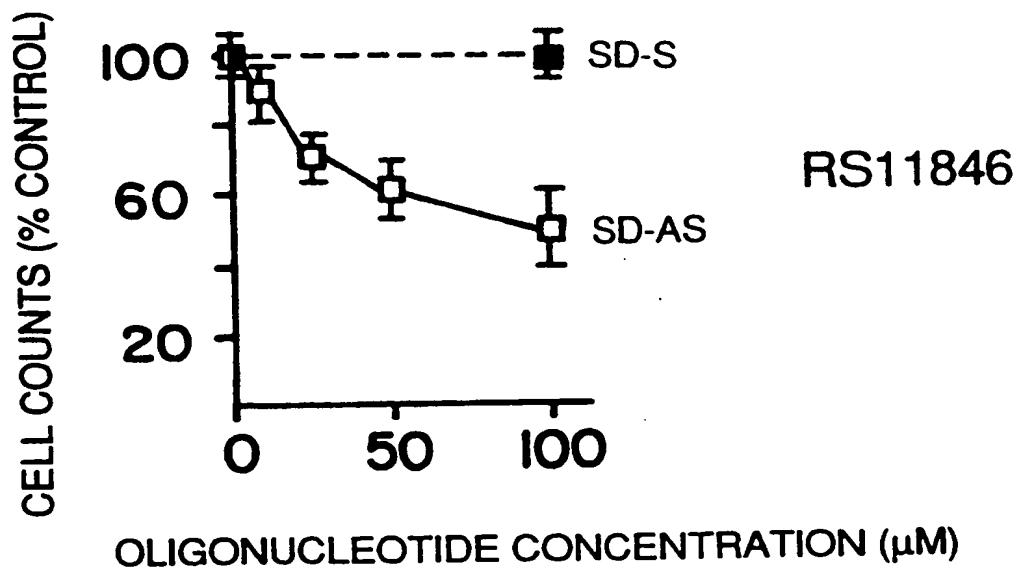


FIG. 1 (b)



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FIG. 1 (c)

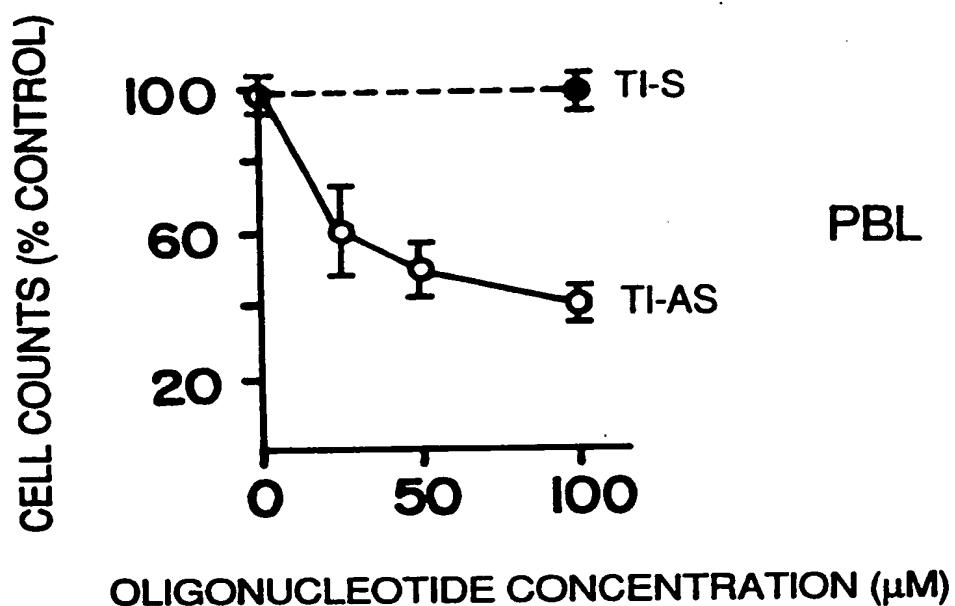
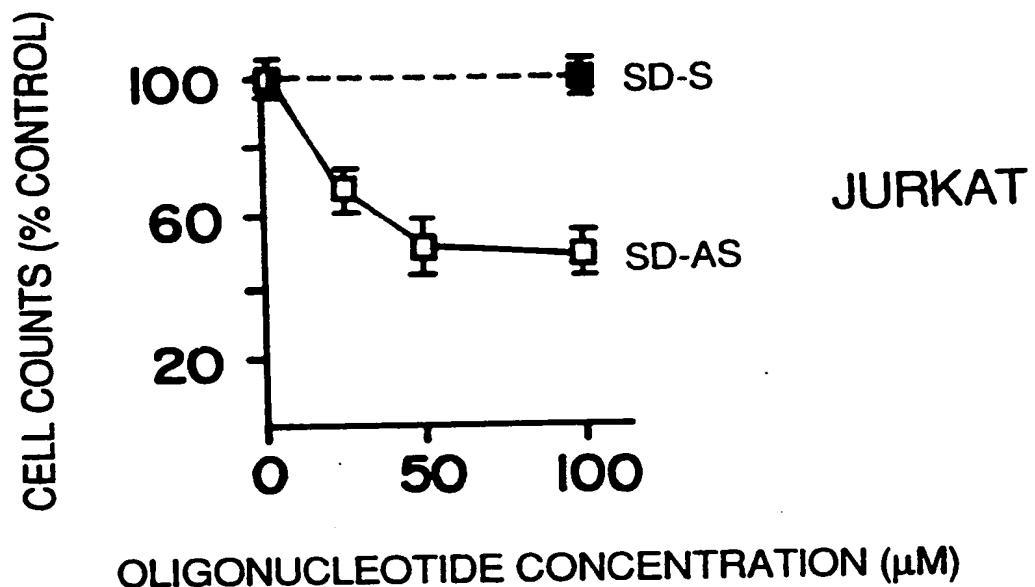


FIG. 1 (d)



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FIG. 2(a)

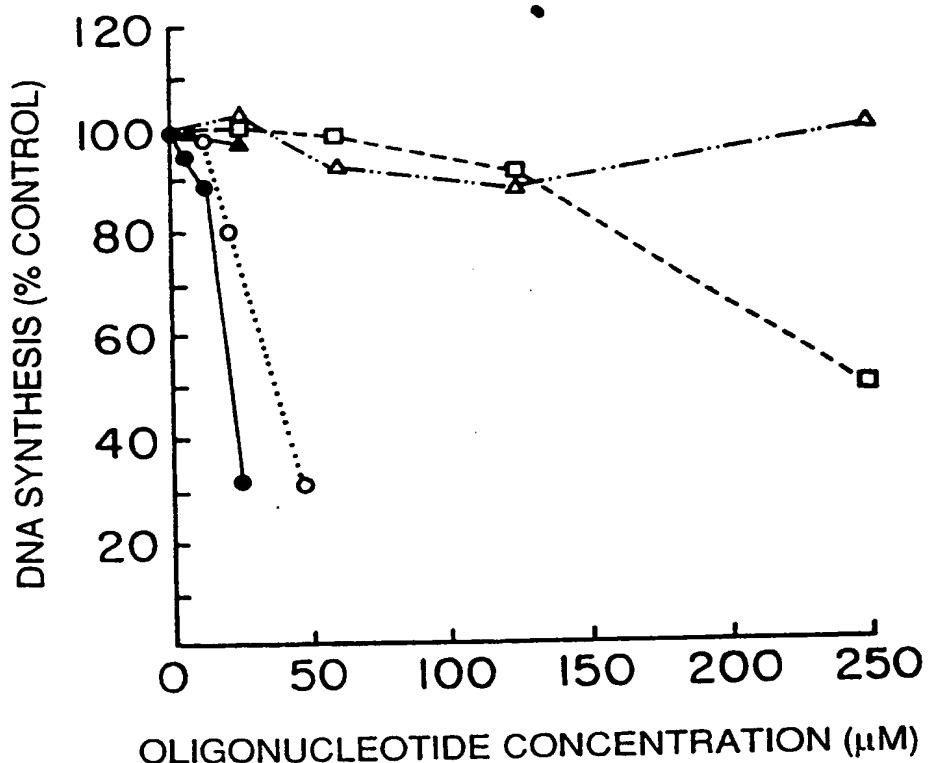
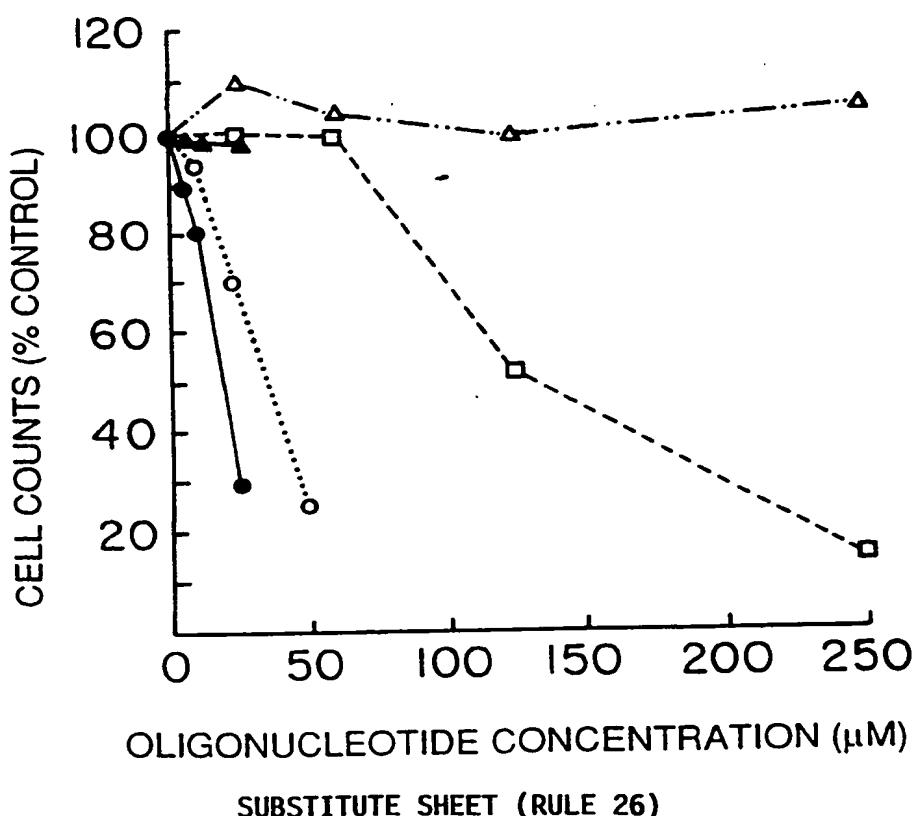


FIG. 2(b)



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011go-2
C-011go 2
011go-1
011go-3
011go-4
011go-5
011go-6
U

FIG. 3

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TI-AS TI-S
SC 20
U
011go 2
C-011go
011go 3

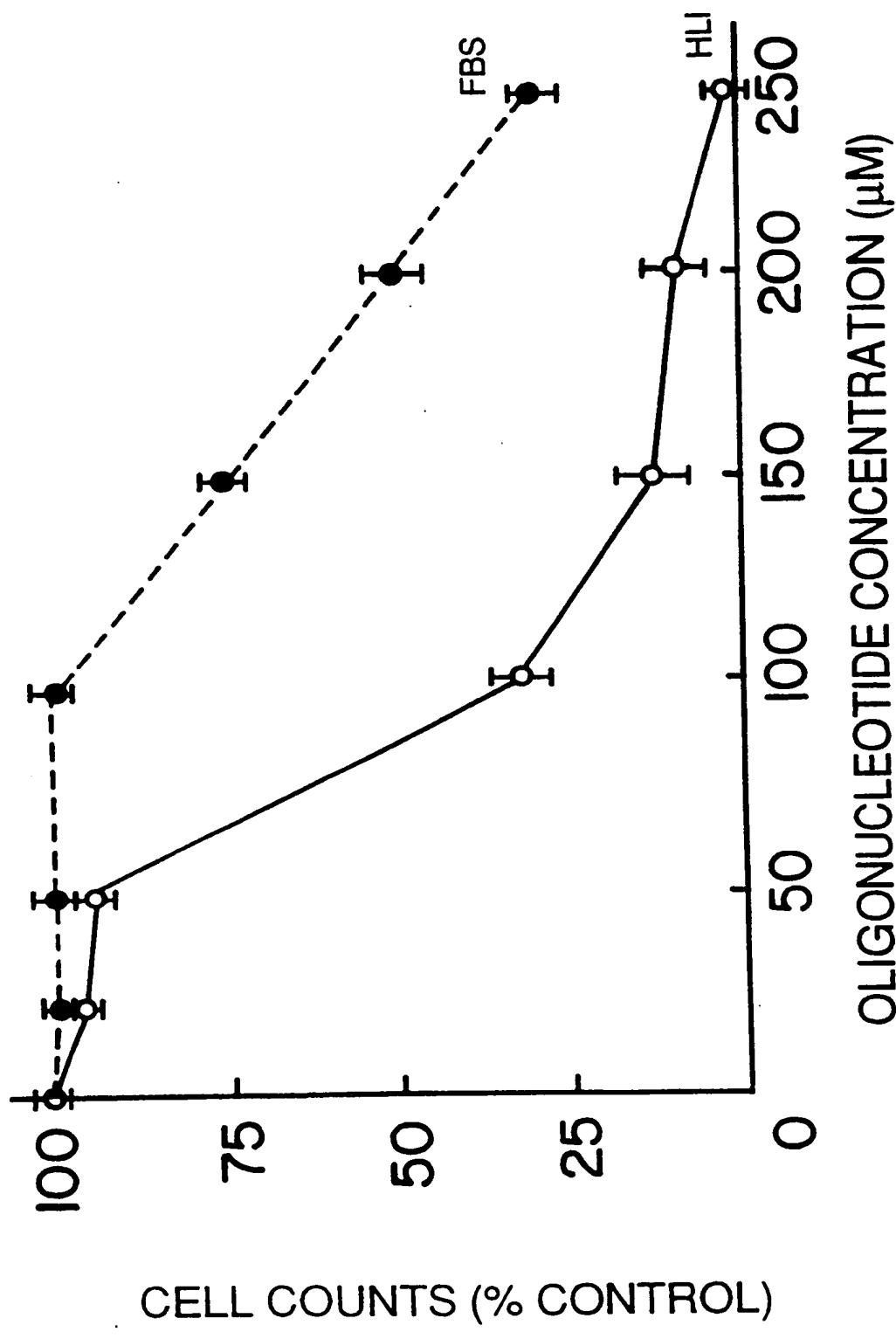
1432
1406
1394
C-011go
U

FIG. 4(a)

FIG. 4(b)

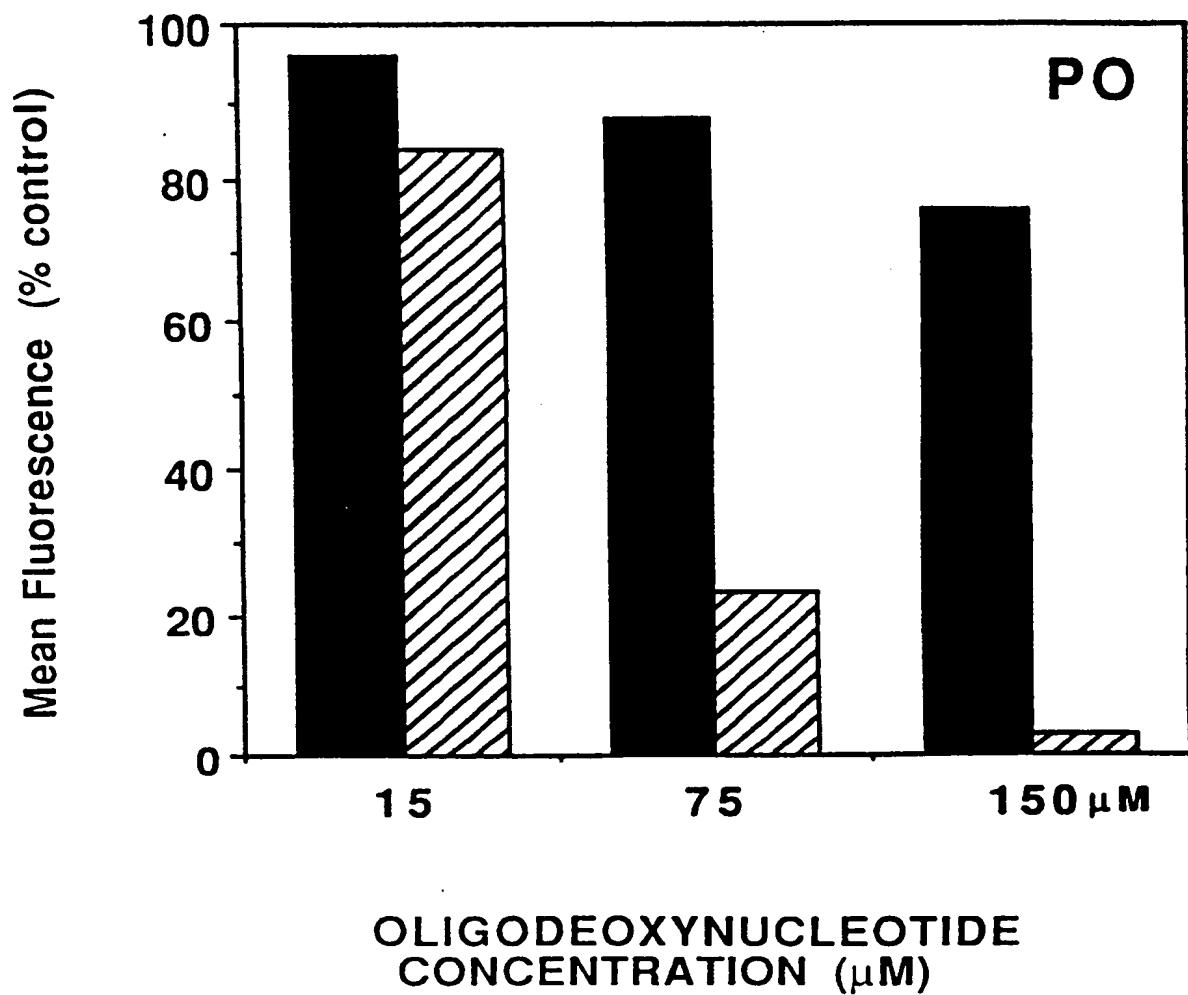
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FIG. 5



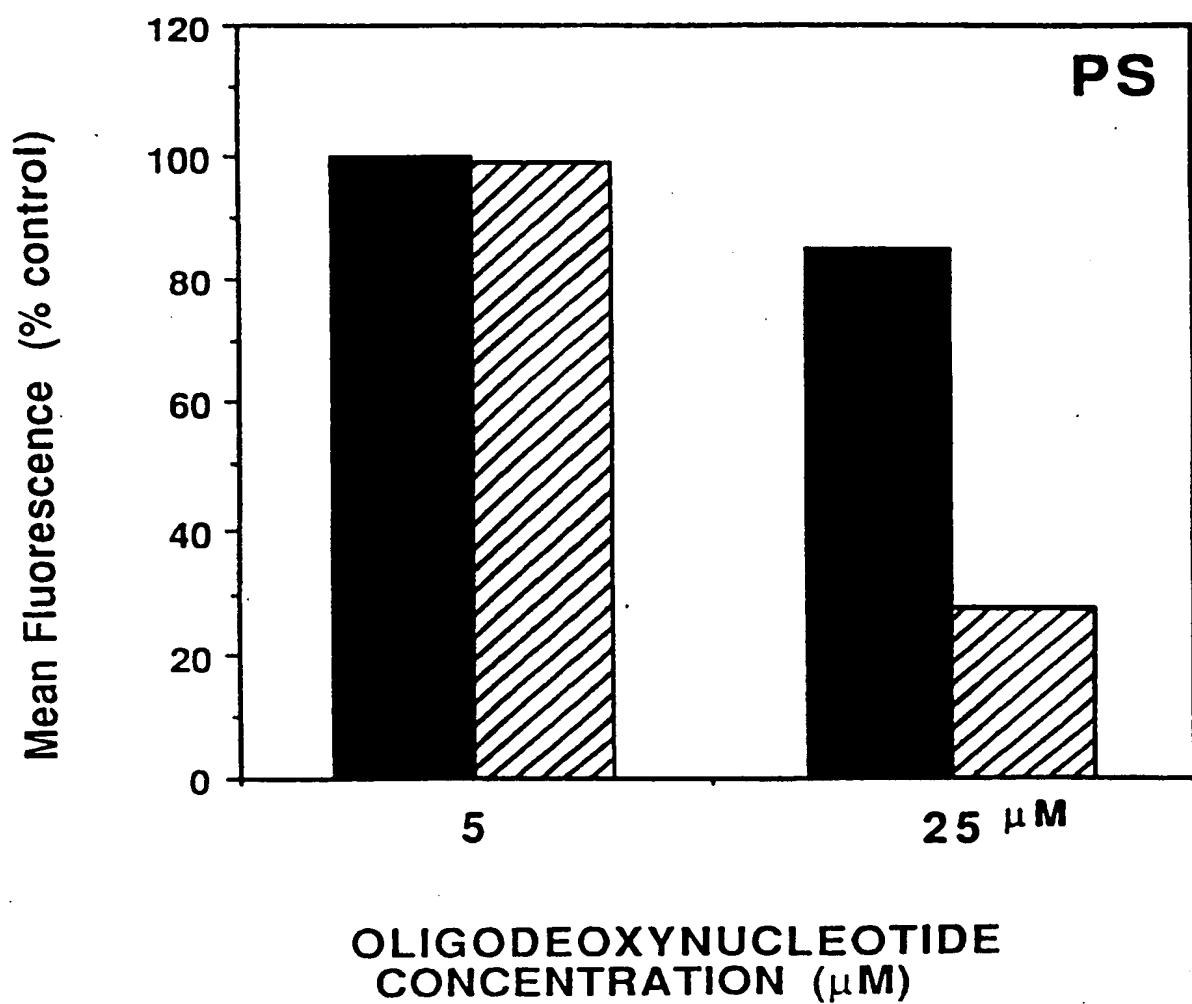
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FIG. 6 (a)



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FIG. 6(b)



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FIG. 7(a)

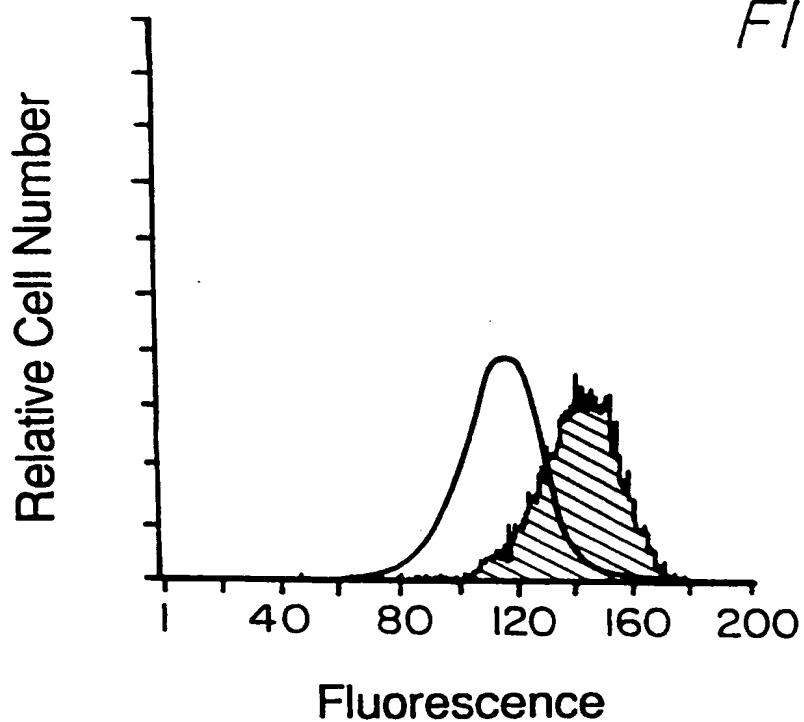
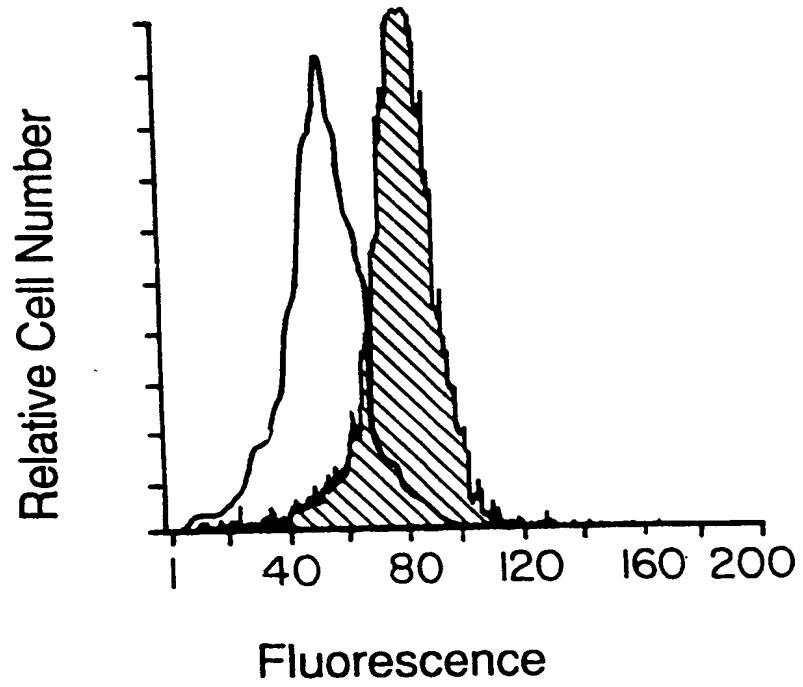


FIG. 7(b)



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FIG. 7(c)

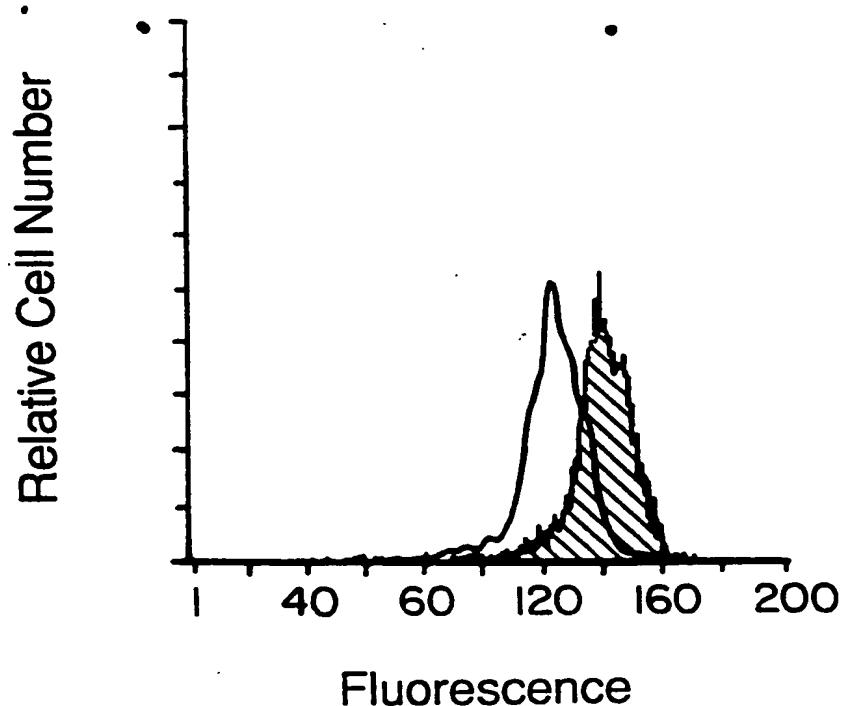
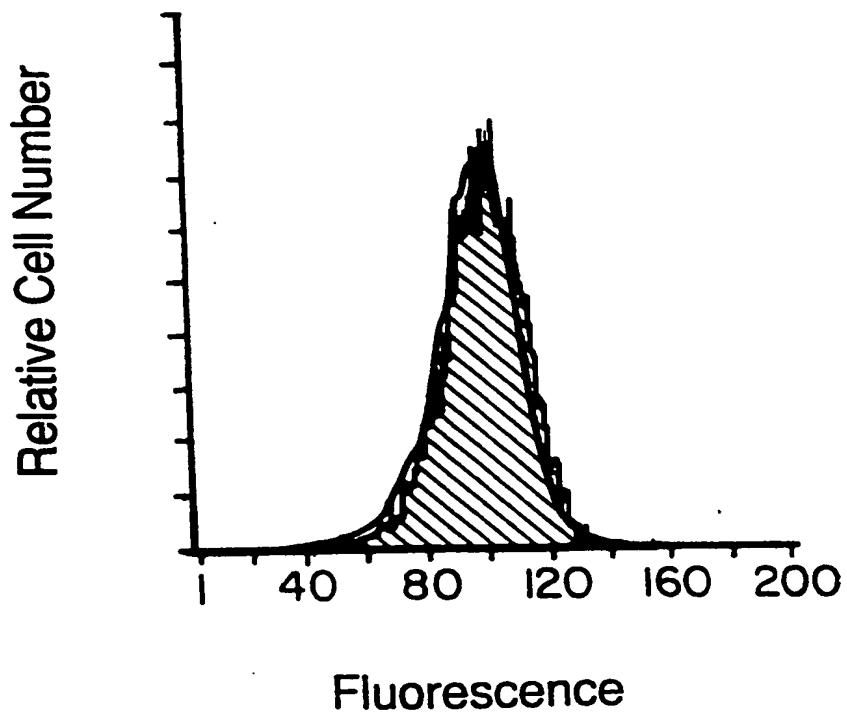
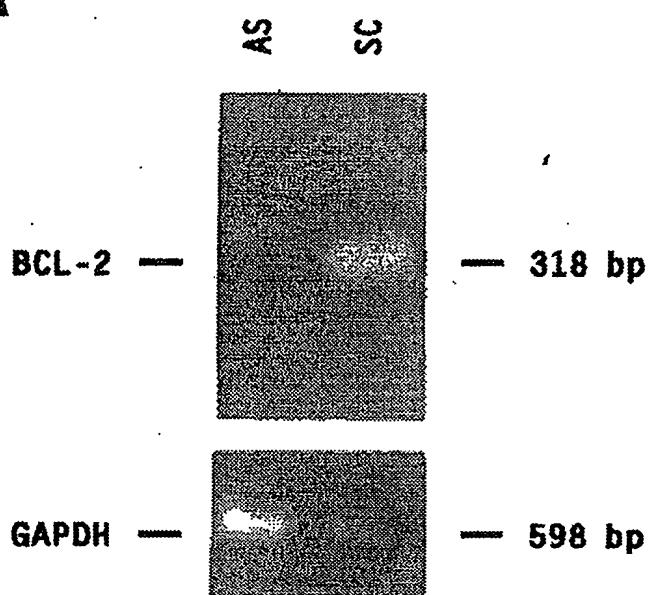
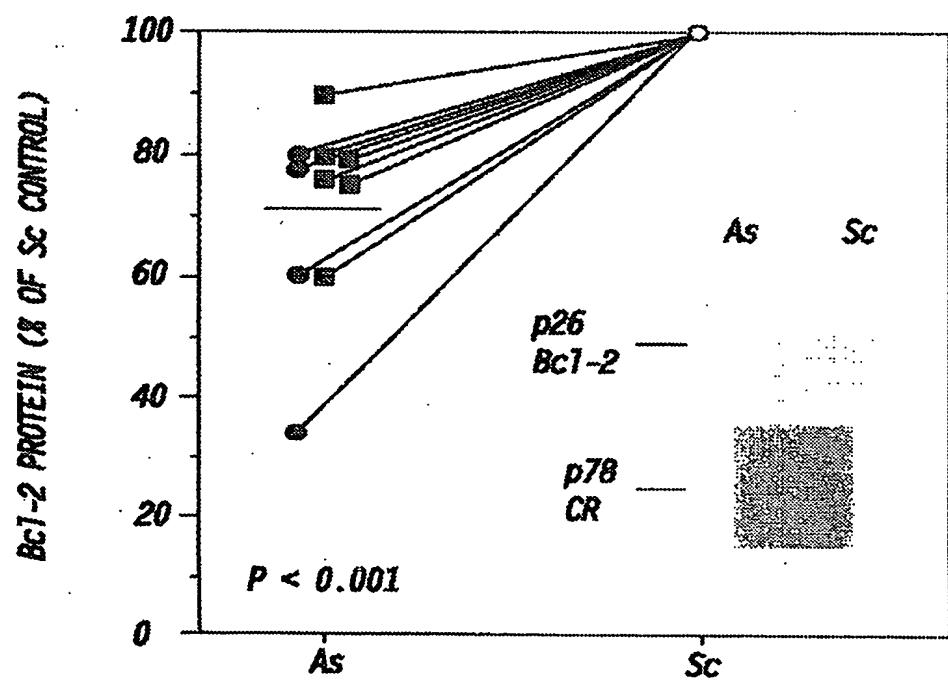


FIG. 7(d)

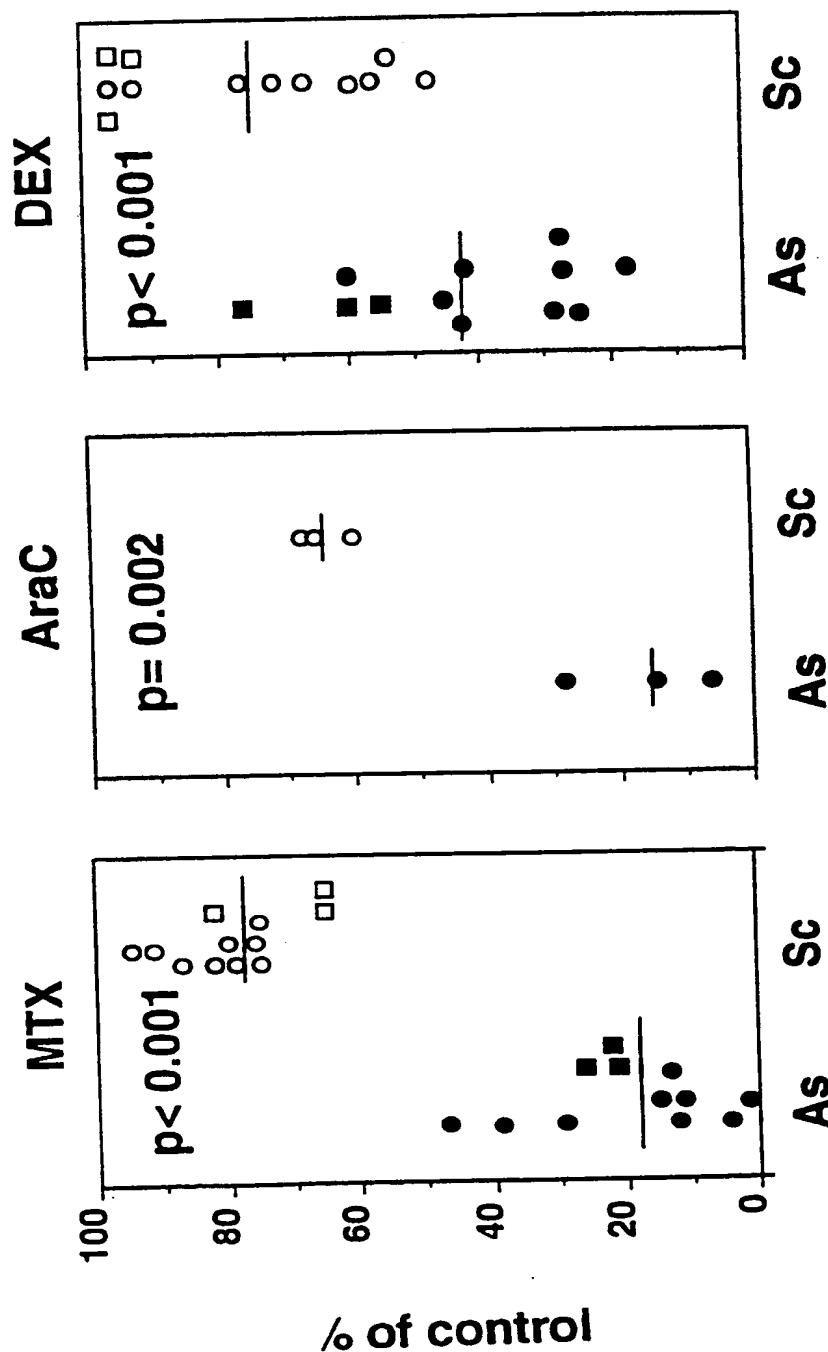


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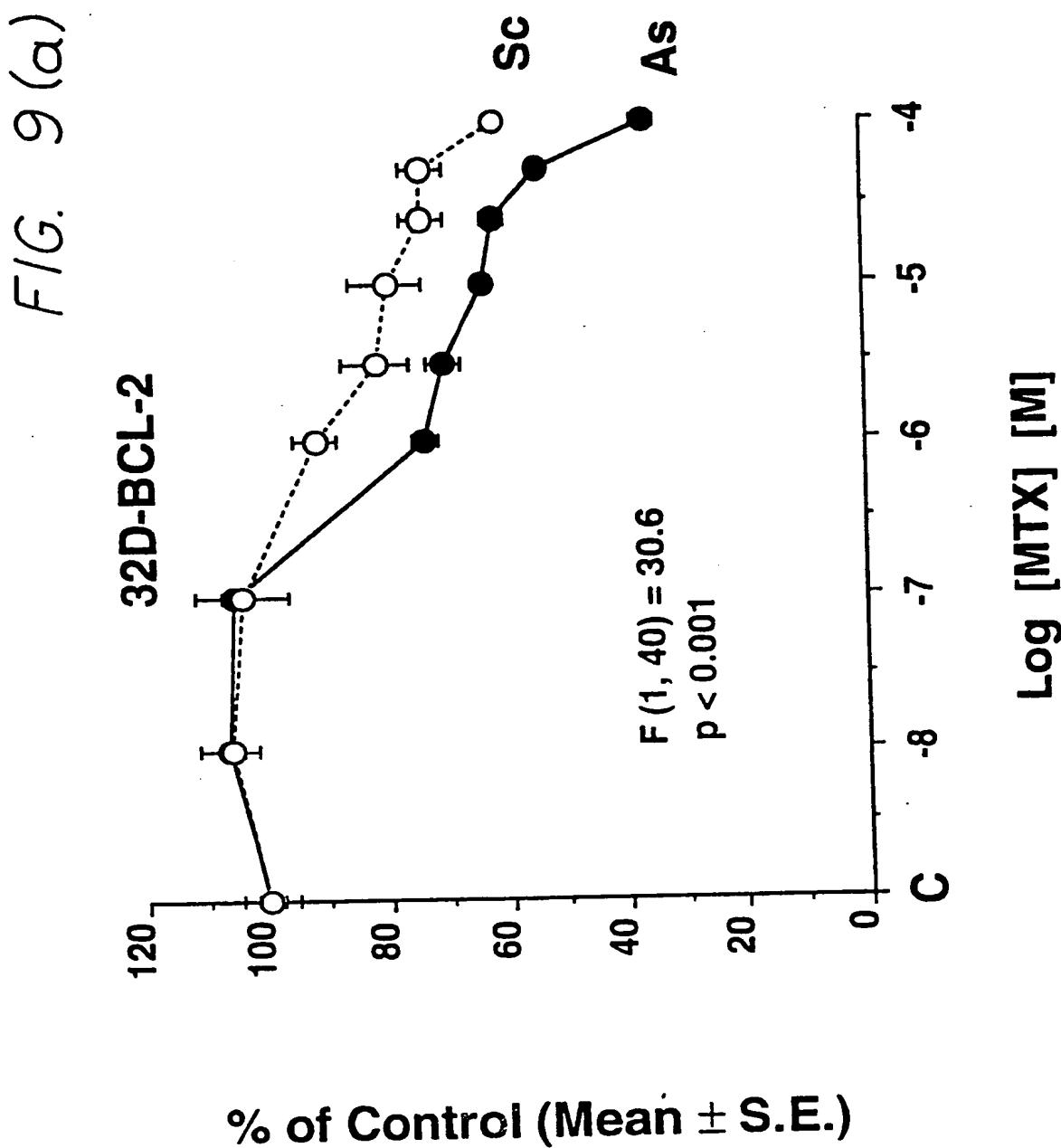
FIG. 8A**FIG. 8B**

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FIG. 8C

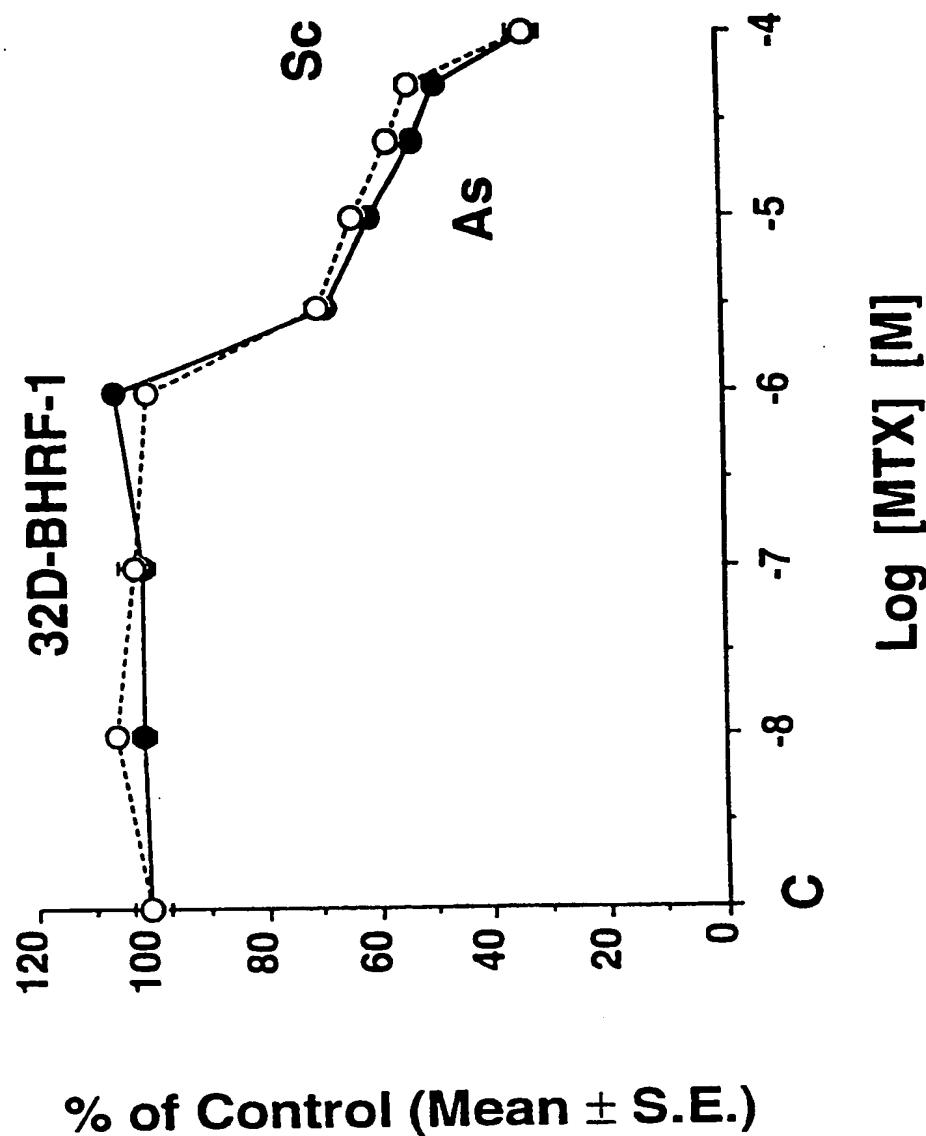


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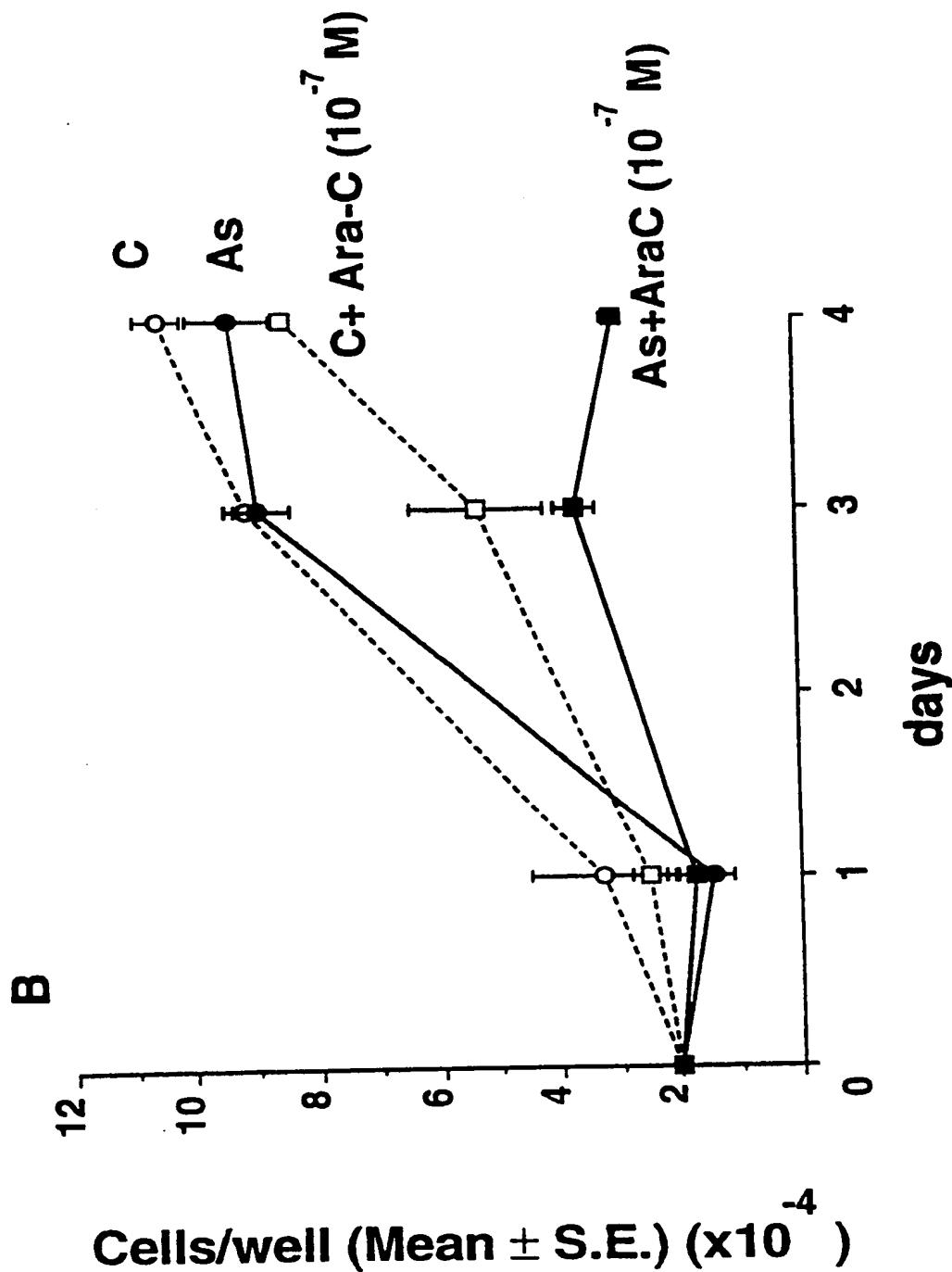
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FIG. 9(b)



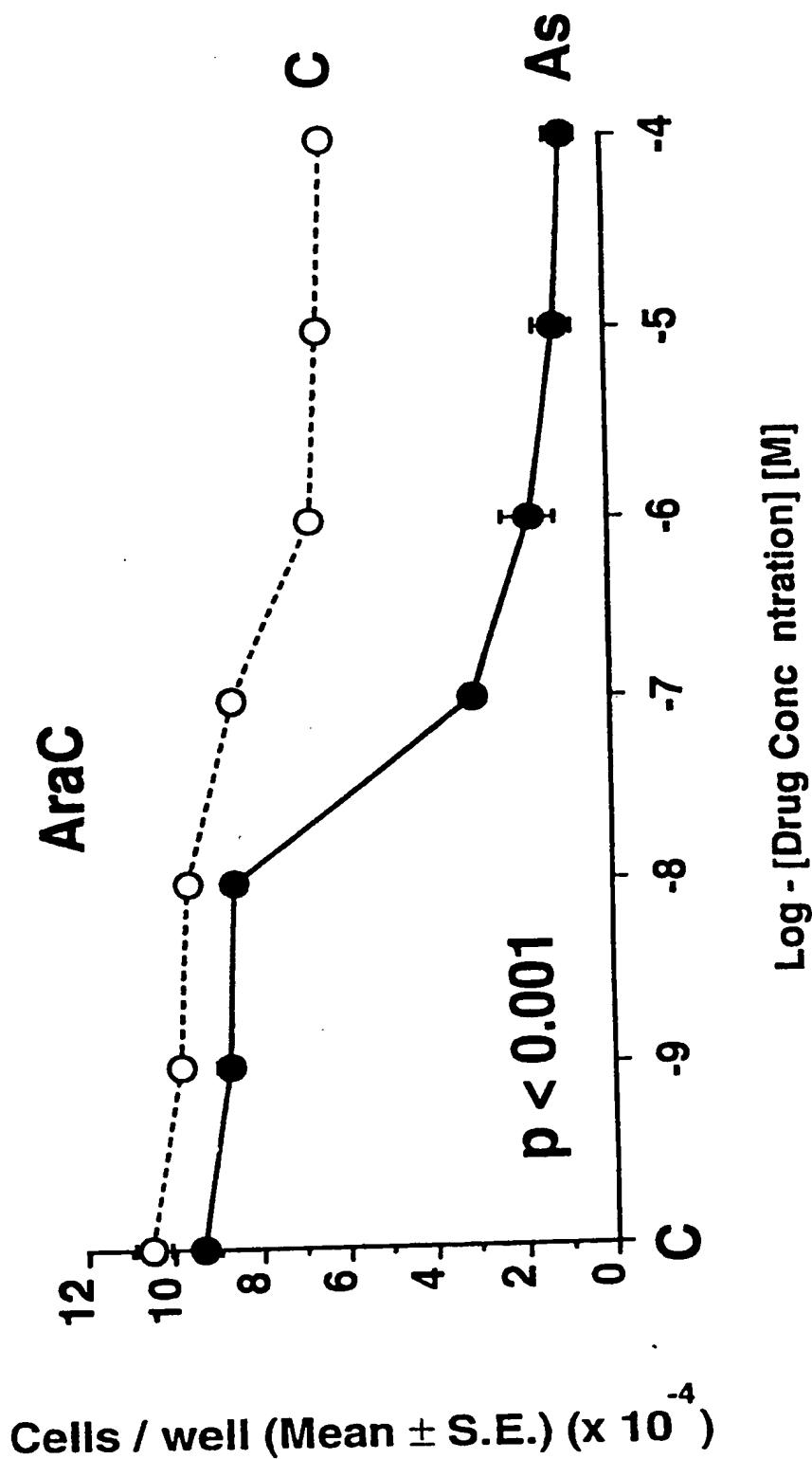
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FIG. 10A



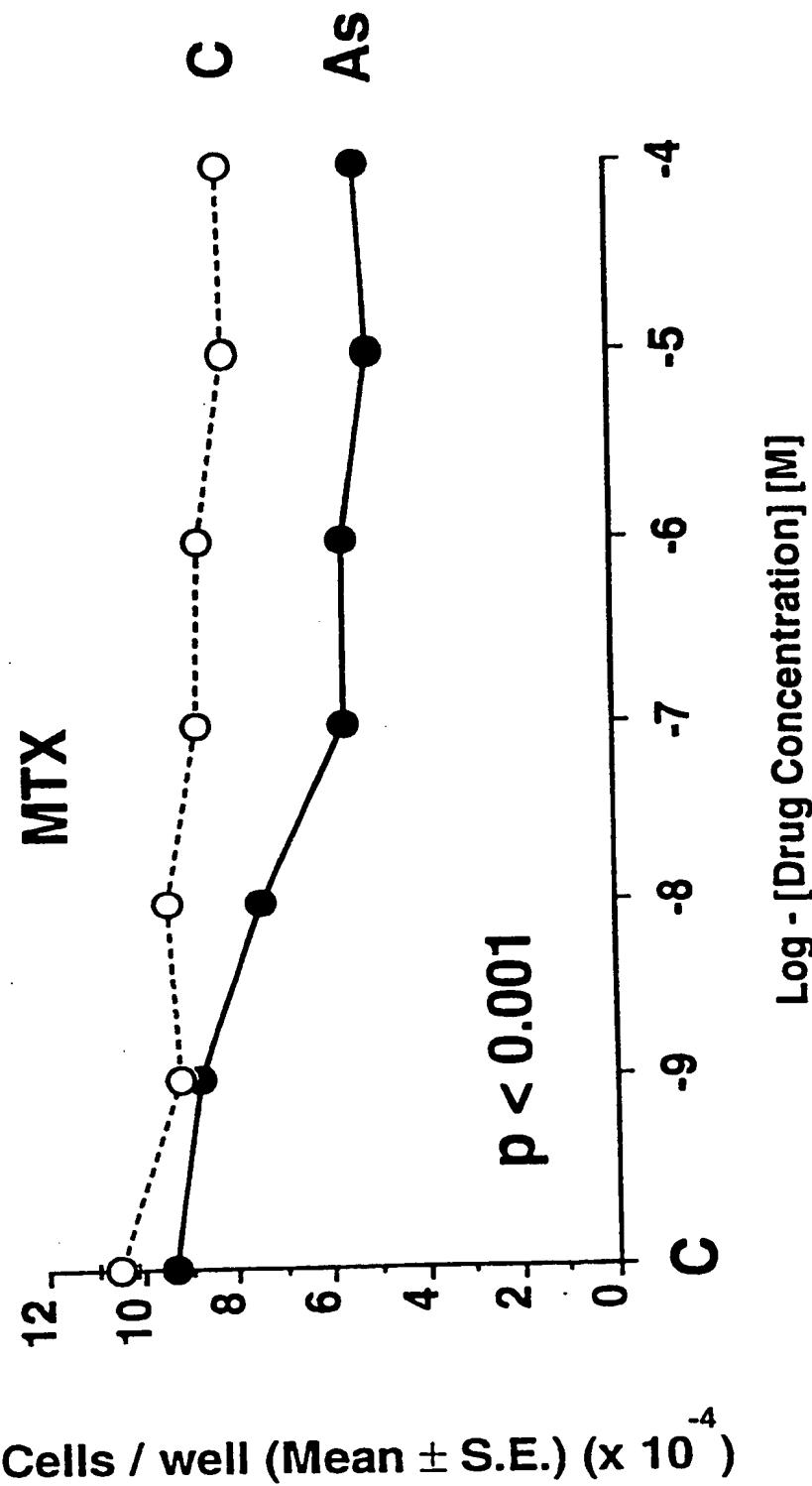
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FIG. 10B (I)



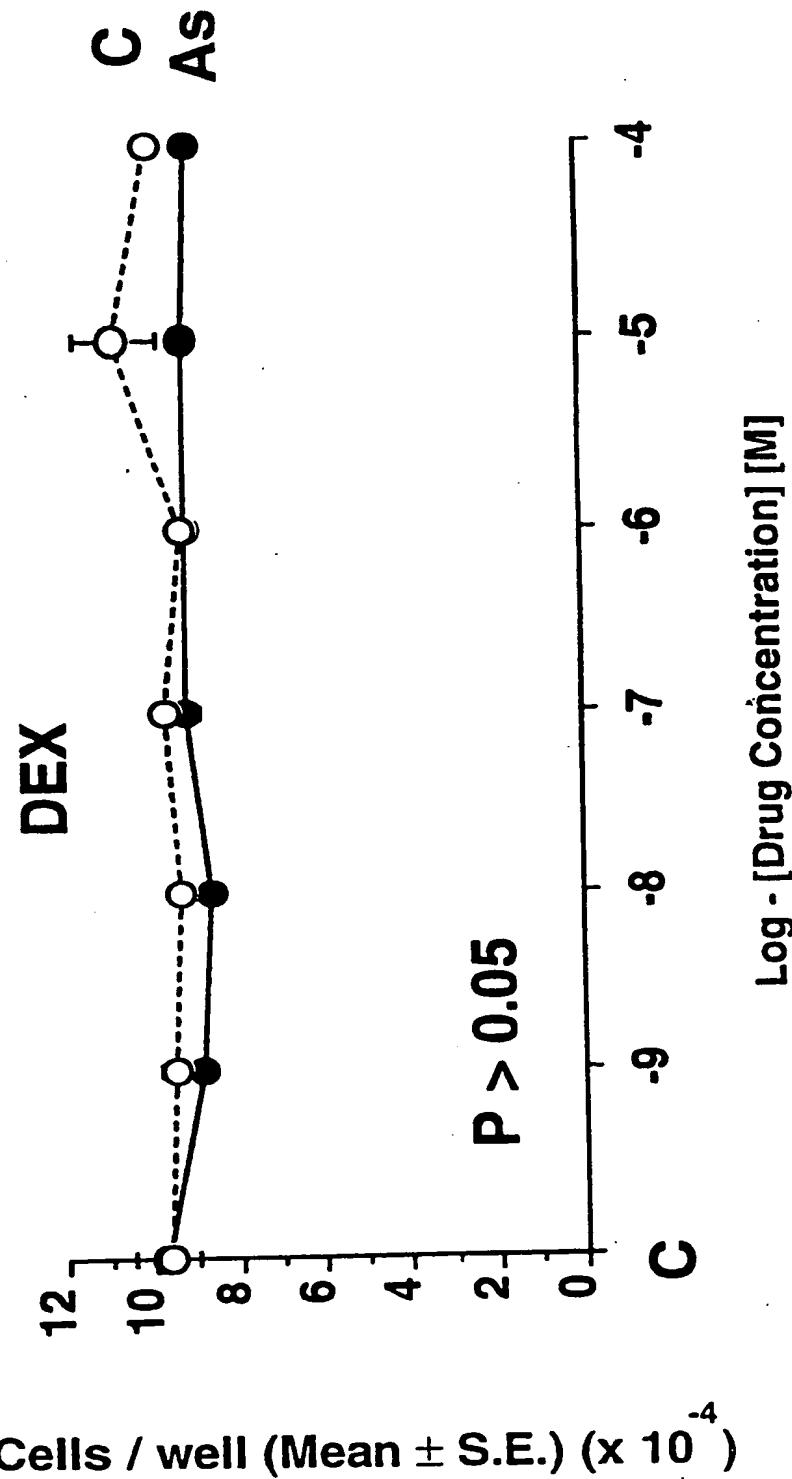
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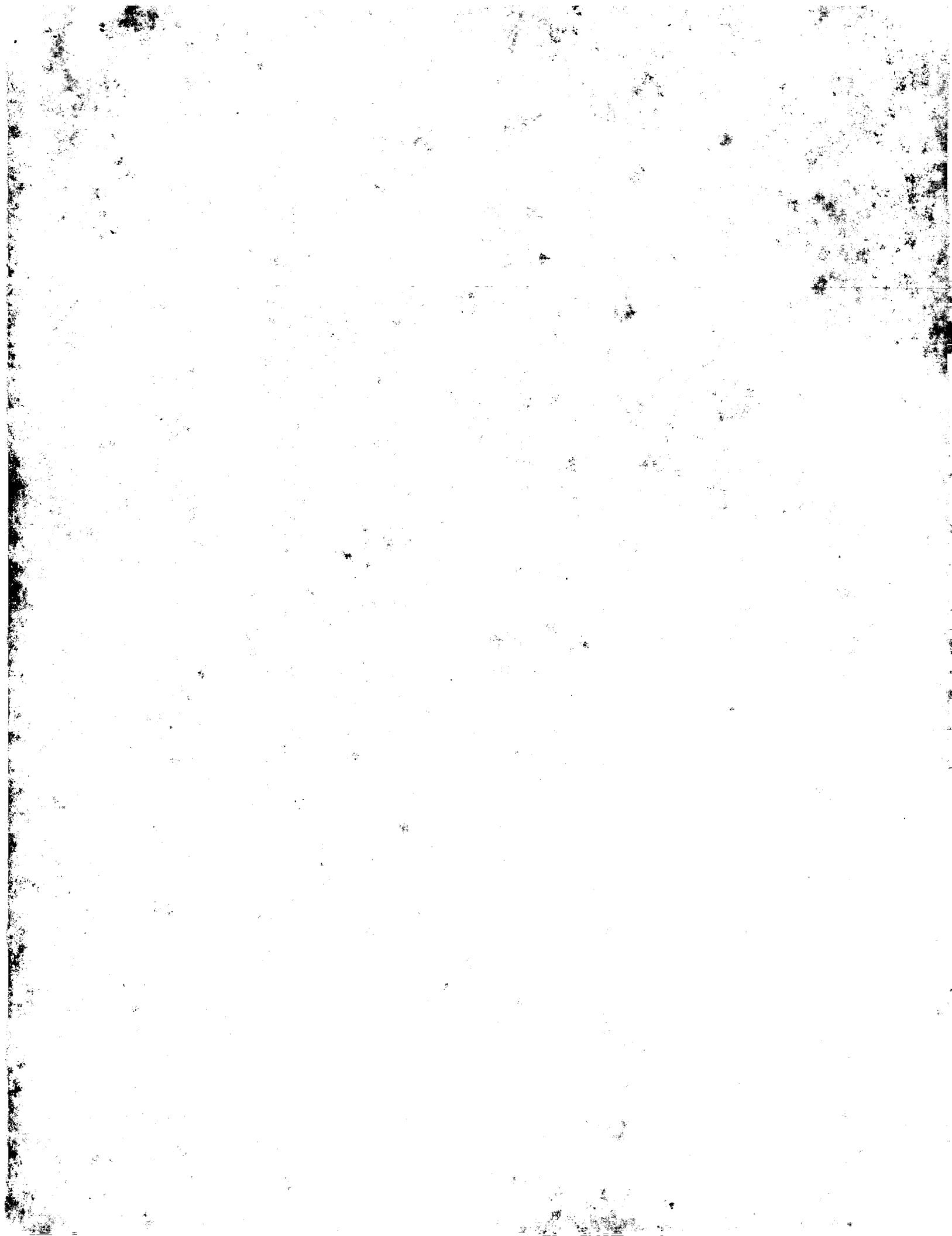
FIG. 10B(2)



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FIG. 10B (3)

Cells / well (Mean ± S.E.) ($\times 10^{-4}$)



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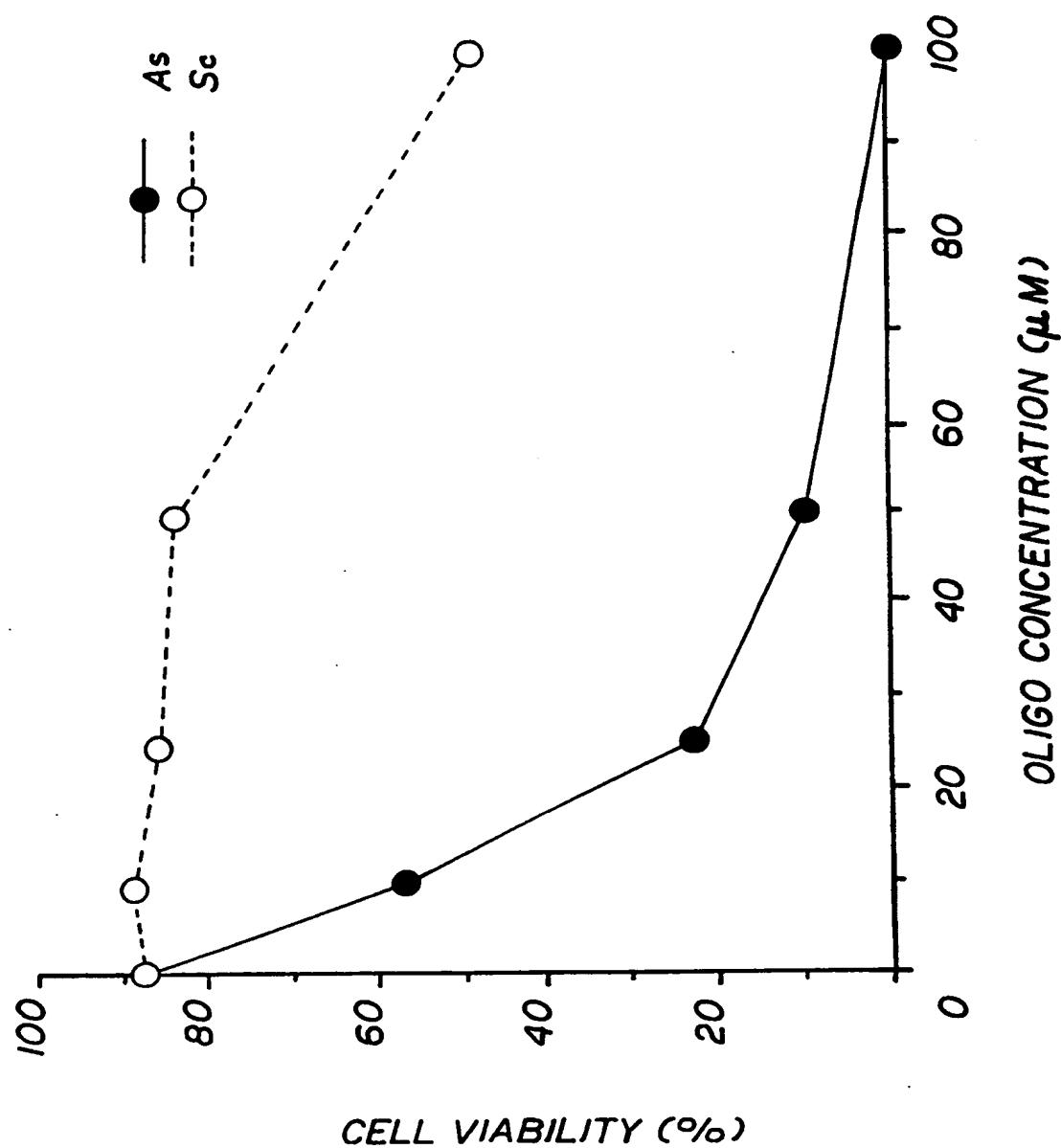


FIG. II

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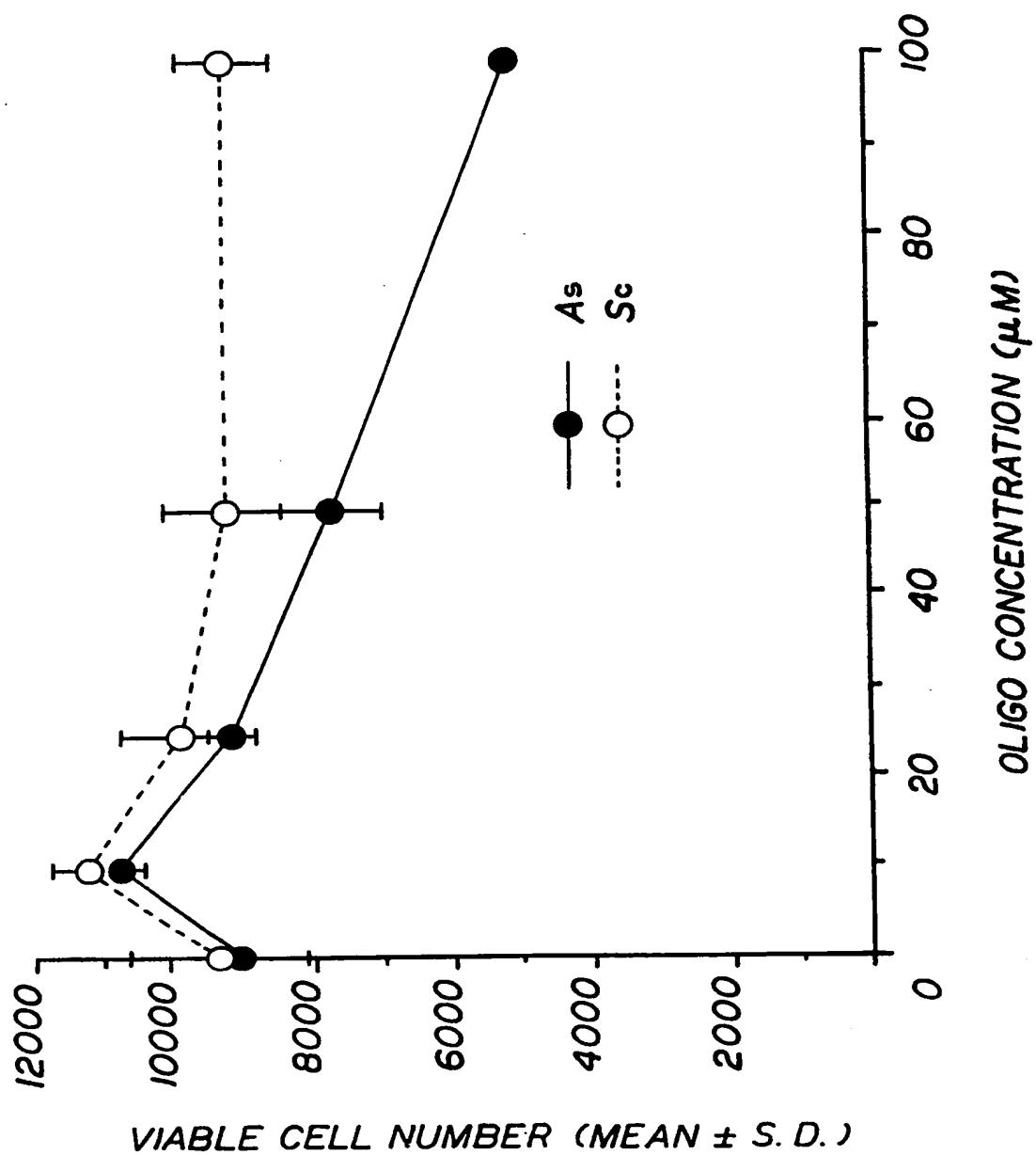
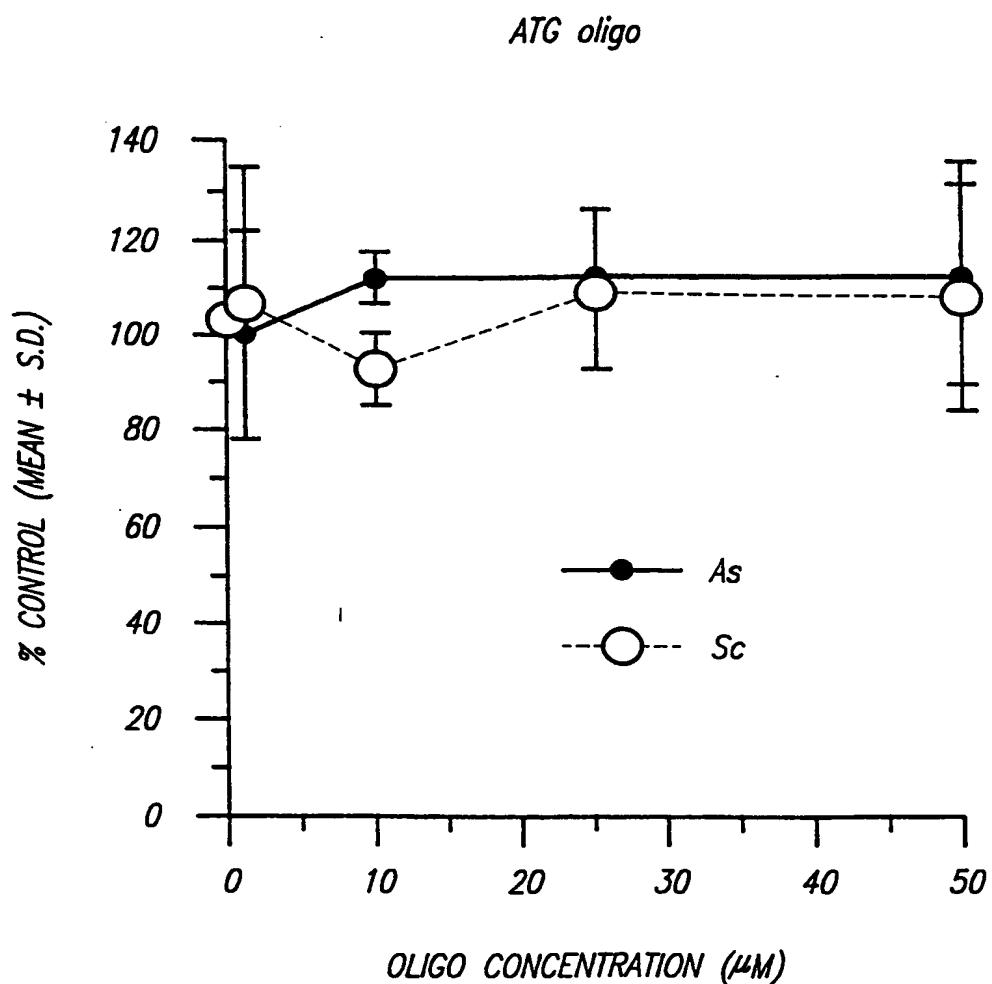


FIG. 12

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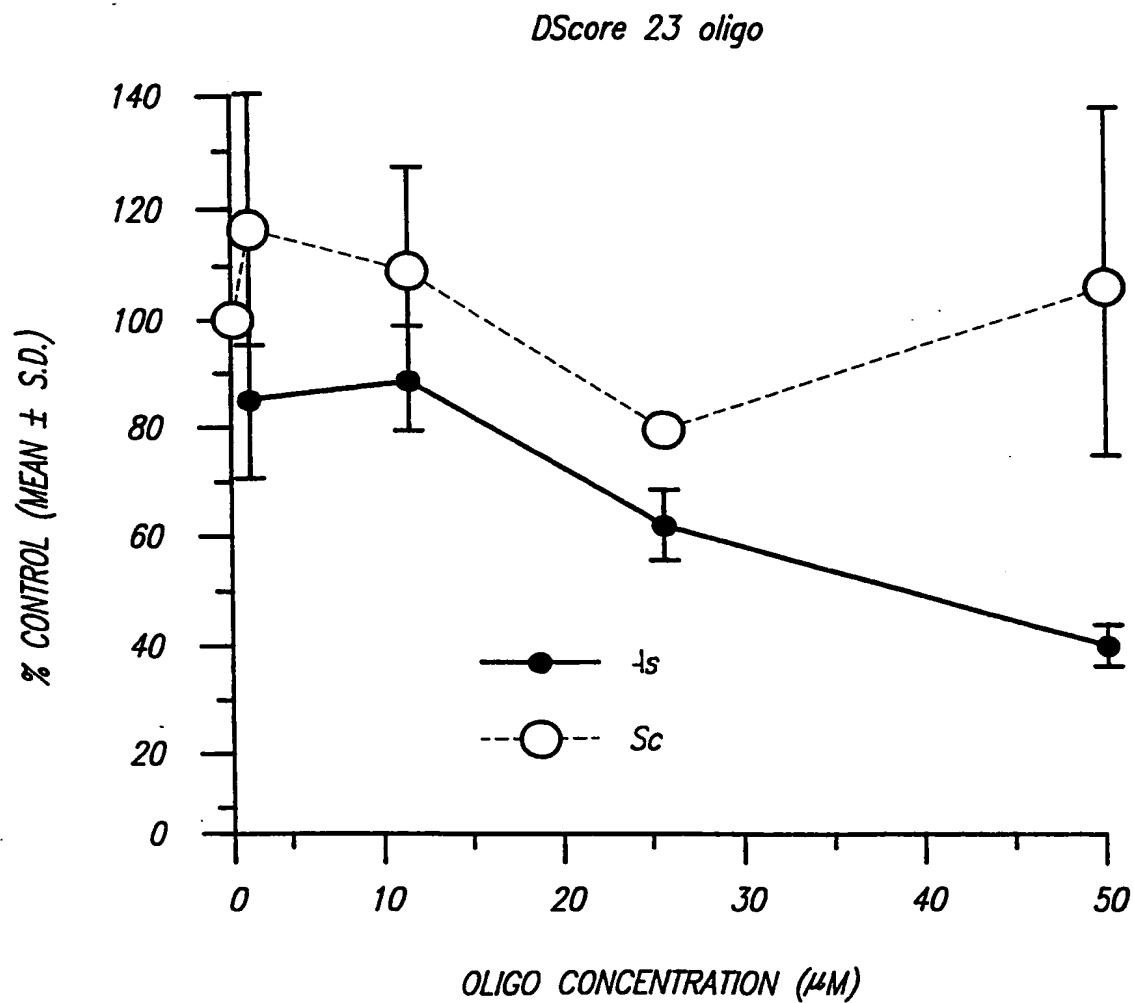
FIG. 13(a)



As-18'mer
5'-TCTCCCAGCGTGCGCCAT-3'
Sc-18'mer
5'-TGCACTCACGCTCGGCCT-3'

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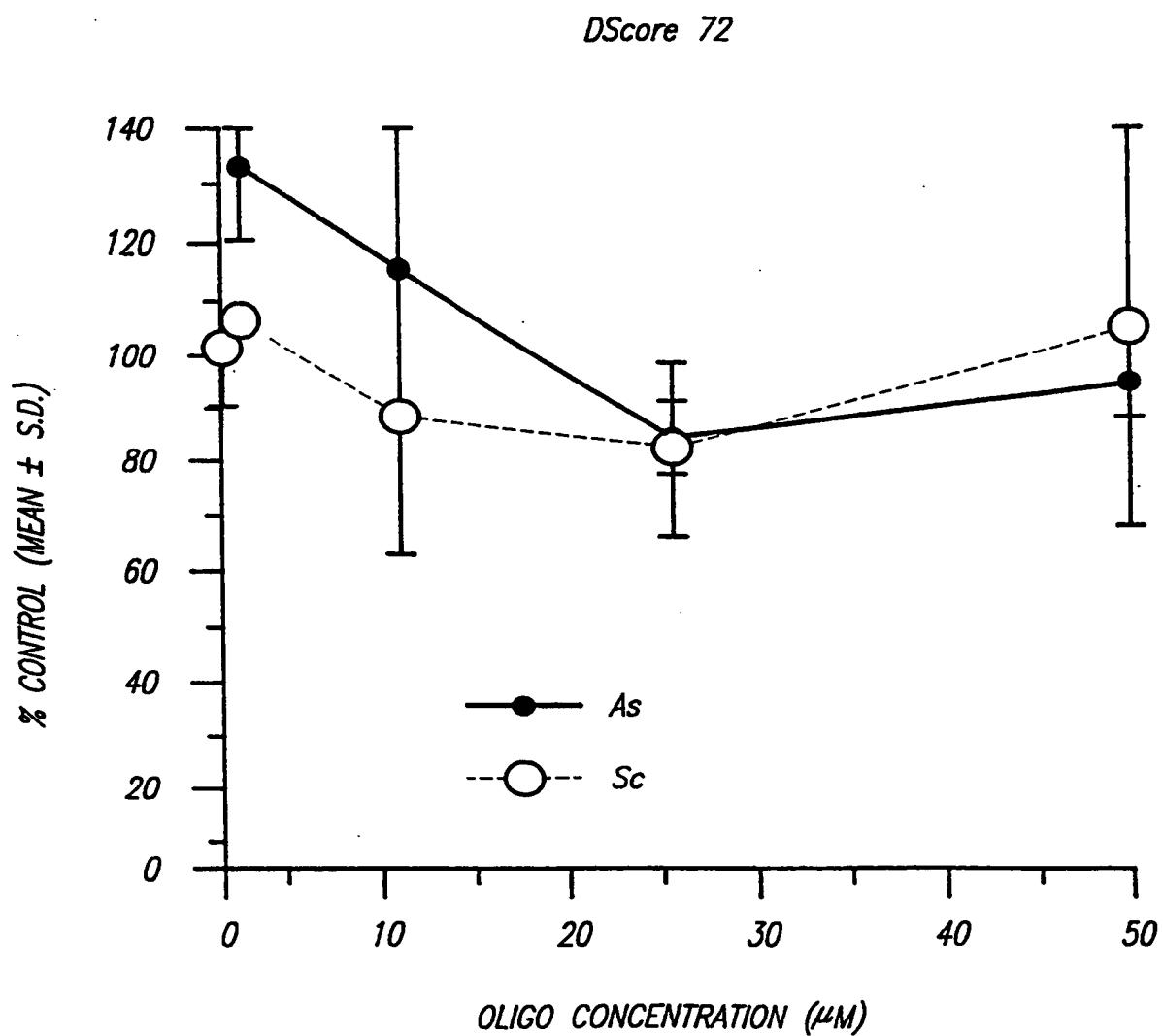
FIG. 13(b)



As23
5'-GCAGGGCGGGCGGGCA-3'
Sc23
5'-GGGCGGAGGCCGGCCGGCGG-3'

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FIG. 13(c)



As72
5'-AGCGGCCGGCGGCCGGCAGCGC-3'
Sc72
5'-GGGCCGGGAAGGGCGCCCCGC-3'

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US94/10725

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : A61K 48/00

US CL : 514/44

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/44

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

None

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS, DIALOG, bcl-2, antisense, anticode, apoptosis, tumor cells, therapeutic agent

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	Anti-Cancer Drug Design, Volume 2, issued 1988, Paoletti, "Anti-sense oligonucleotides as potential antitumour agents: prospective views and preliminary results", pages 325-331, see entire article.	1-40
Y	Proceedings of the National Academy of Sciences USA, Volume 83, issued July 1986, Tsujimoto et al., "Analysis of the structure, transcripts and protein products of <i>bcl-2</i> , the gene involved in human follicular lymphoma", pages 5214-5218, see entire article.	1-40
Y	Cancer Research, Volume 48, issued 15 May 1988, Stein et al., "Oligodeoxynucleotides as inhibitors of gene expression: A review", pages 2659-2668, see entire article.	1-40



Further documents are listed in the continuation of Box C.



See patent family annex.

• Special categories of cited documents:			
"A"	document defining the general state of the art which is not considered to be of particular relevance	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E"	earlier document published on or after the international filing date	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L"	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O"	document referring to an oral disclosure, use, exhibition or other means	"&"	document member of the same patent family
"P"	document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

08 NOVEMBER 1994

Date of mailing of the international search report

12 JAN 1995

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Facsimil No. (703) 305-3230

Authorized Officer

SUZANNE ZISKA, PH.D.

Telephone No. (703) 308-0196

R. Kuzza Jr.

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	Cancer Research, Volume 50, issued 15 October 1990, Reed et al., "Antisense-mediated inhibition of <i>BCL2</i> protooncogene expression and leukemic cell growth and survival: Comparisons of phosphodiester and phosphorothioate oligodeoxynucleotides", pages 6565-6570, see entire article.	1-40
Y	Cell, Volume 67, issued 29 November 1991, Strasser et al., " <i>bcl-2</i> transgene inhibits T cell death and perturbs thymic self-censorship", pages 889-899, see entire article.	1-40
Y	Chemical Reviews, Volume 90, No. 4, issued June 1990, Uhlmann et al., "Antisense oligonucleotides: A new therapeutic principle", pages 544-584, see entire article.	1-40
Y	US, A, 5,098,890 (GEWIRTZ ET AL.) 24 March 1992, see entire document.	1-40